



(12) **EUROPEAN PATENT APPLICATION**
published in accordance with Art. 153(4) EPC

(43) Date of publication:
09.02.2022 Bulletin 2022/06

(21) Application number: **21780843.5**

(22) Date of filing: **08.02.2021**

(51) International Patent Classification (IPC):
C08L 13/02 ^(2006.01) **C08L 9/04** ^(2006.01)
C08J 3/02 ^(2006.01) **C08F 236/12** ^(2006.01)
C08K 3/011 ^(2018.01) **C08K 3/22** ^(2006.01)
C08J 5/02 ^(2006.01) **B29C 41/14** ^(2006.01)

(52) Cooperative Patent Classification (CPC):
B29C 41/14; C08F 236/12; C08J 3/02; C08J 5/02;
C08K 3/011; C08K 3/22; C08L 9/04; C08L 13/02

(86) International application number:
PCT/KR2021/001626

(87) International publication number:
WO 2021/201416 (07.10.2021 Gazette 2021/40)

(84) Designated Contracting States:
AL AT BE BG CH CY CZ DE DK EE ES FI FR GB
GR HR HU IE IS IT LI LT LU LV MC MK MT NL NO
PL PT RO RS SE SI SK SM TR
Designated Extension States:
BA ME
Designated Validation States:
KH MA MD TN

(30) Priority: **31.03.2020 KR 20200039060**
31.03.2020 KR 20200039065
31.03.2020 KR 20200039067
31.03.2020 KR 20200039071
31.03.2020 KR 20200039074
31.03.2020 KR 20200039076
11.12.2020 KR 20200172992

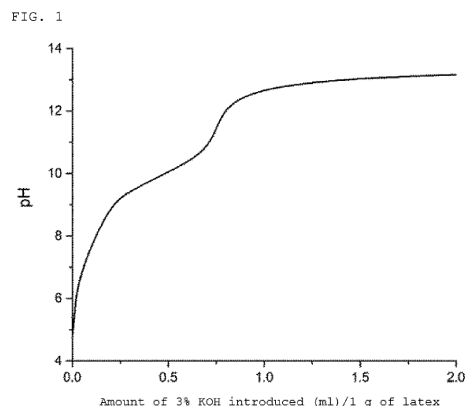
(71) Applicant: **LG CHEM, LTD.**
Yeongdeungpo-gu,
Seoul 07336 (KR)

(72) Inventors:
• **OH, Seung Whan**
Daejeon 34122 (KR)
• **KWON, Won Sang**
Daejeon 34122 (KR)
• **KIM, Jung Eun**
Daejeon 34122 (KR)
• **KIM, Ji Hyun**
Daejeon 34122 (KR)
• **YEU, Seung Uk**
Daejeon 34122 (KR)
• **JANG, Myung Su**
Daejeon 34122 (KR)
• **HA, Do Young**
Daejeon 34122 (KR)
• **HAN, Jung Su**
Daejeon 34122 (KR)

(74) Representative: **Plasseraud IP**
66, rue de la Chaussée d'Antin
75440 Paris Cedex 09 (FR)

(54) **CARBOXYLIC ACID-MODIFIED NITRILE-BASED COPOLYMER LATEX, LATEX COMPOSITION FOR DIP MOLDING COMPRISING SAME, AND MOLDED ARTICLE FORMED THEREFROM**

(57) The present invention relates to a latex composition for dip molding comprising a carboxylic acid-modified nitrile-based copolymer latex satisfying General Formulas 1 and 2 (see the specification), wherein the carboxylic acid-modified nitrile-based copolymer includes a conjugated diene-based monomer-derived unit, an ethylenic unsaturated nitrile-based monomer-derived unit, and an ethylenic unsaturated acid monomer-derived unit, and a molded article.



Description

[Technical Field]

CROSS-REFERENCE TO RELATED APPLICATIONS

[0001] The present application claims the benefit of priorities to Korean Patent Application No. 10-2020-0039060, filed on March 31, 2020, Korean Patent Application No. 10-2020-0039067, filed on March 31, 2020, Korean Patent Application No. 10-2020-0039076, filed on March 31, 2020, Korean Patent Application No. 10-2020-0039074, filed on March 31, 2020, Korean Patent Application No. 10-2020-0039065, filed on March 31, 2020, Korean Patent Application No. 10-2020-0039071, filed on March 31, 2020, and Korean Patent Application No. 10-2020-0172992, filed on December 11, 2020, the entire contents of which are incorporated herein by reference.

Technical Field

[0002] The present invention relates to a carboxylic acid-modified nitrile-based copolymer latex capable of producing a dip-molded article having high elasticity and softness, a latex composition for dip molding comprising the copolymer latex, and a molded article formed of the latex composition.

[Background Art]

[0003] Usage of a disposable rubber glove has increased in various fields such as housework, the food industry, the electronic industry, and medical fields. In the past, a disposable glove was made by dip molding a natural rubber latex, but a protein was problematic due to an allergic reaction, such as pain or a rash, in some users. Due to these problems, recently, a disposable glove formed by dip molding a nitrile-based latex, which has a high tensile strength, an excellent chemical resistance, and no risk of allergy, has been spotlighted in order to replace natural rubber.

[0004] However, since the nitrile-based rubber glove obtained by using the nitrile-based latex is inferior to a natural rubber glove in terms of elasticity and softness, the nitrile-based rubber glove still cannot replace the natural rubber glove in uses such as a surgical glove where excellent wearing comfort and maintenance of adhesion when wearing for a long time are important.

[0005] Therefore, various studies have been conducted to improve both the softness and elasticity of the glove. However, conventionally, studies on a change of a cross-linking agent such as changing a ratio between zinc oxide, sulfur, and a vulcanization accelerator to be added together with the latex in the production of the glove, or adding a new cross-linking agent have been mainly conducted rather than studies on a latex required for producing a glove having high elasticity and excellent wearing comfort. However, in a case where the elasticity and softness are improved through only the change of the cross-linking agent, other physical properties, such as a tensile strength and durability, of the glove may be deteriorated. In addition, in a case where sulfur or a vulcanization accelerator is excessively used, discoloration of the glove or an IV type allergy may occur.

[0006] Therefore, there is a need for development of a latex itself to improve both the elasticity and softness of the nitrile-based glove.

[Disclosure]**[Technical Problem]**

[0007] In order to solve the problems mentioned in the background art, an object of the present invention is to provide a latex composition for dip molding comprising a carboxylic acid-modified nitrile-based copolymer latex to improve elasticity and softness of a nitrile-based rubber glove.

[0008] That is, an object of the present invention is to improve elasticity and softness of a nitrile-based rubber glove formed of a latex composition for dip molding comprising a carboxylic acid-modified nitrile-based copolymer latex by adjusting a capillary viscosity (CV_0) of the carboxylic acid-modified nitrile-based copolymer latex in a swollen state in a methyl ethyl ketone solvent to a range of 1.0 mm²/s to 3.0 mm²/s, and adjusting a ratio P of a capillary viscosity (CV_D) of the carboxylic acid-modified nitrile-based copolymer latex in a de-swollen state to the capillary viscosity (CV_0) of the carboxylic acid-modified nitrile-based copolymer latex in the swollen state in the methyl ethyl ketone solvent, to a range of 0.85 to 1.0.

[Technical Solution]

[0009] In one general aspect, a latex composition for dip molding comprises a carboxylic acid-modified nitrile-based copolymer latex satisfying the following General Formula 1 and General Formula 2, wherein the carboxylic acid-modified nitrile-based copolymer includes a conjugated diene-based monomer-derived unit, an ethylenic unsaturated nitrile-based monomer-derived unit, and an ethylenic unsaturated acid monomer-derived unit:

[General Formula 1]

$$1.0 \text{ mm}^2/\text{s} \leq CV_0 \leq 3.0 \text{ mm}^2/\text{s}$$

[General Formula 2]

$$0.85 \leq P \leq 1.0, P = CV_D/CV_0$$

in General Formulas 1 and 2, CV_0 represents a capillary viscosity of the carboxylic acid-modified nitrile-based copolymer latex measured in a swollen state in a methyl ethyl ketone solvent, and CV_D represents a capillary viscosity of the carboxylic acid-modified nitrile-based copolymer latex in a de-swollen state.

[0010] In another general aspect, a latex composition for dip molding comprises the carboxylic acid-modified nitrile-based copolymer latex and a cross-linking agent composition.

[0011] In still another general aspect, a molded article includes the latex composition for dip molding-derived layer.

[Advantageous Effects]

[0012] According to the carboxylic acid-modified nitrile-based copolymer latex according to the present invention, the capillary viscosity (CV_0) of the carboxylic acid-modified nitrile-based copolymer latex in the swollen state in the methyl ethyl ketone solvent is adjusted to a range of 1.0 mm²/s to 3.0 mm²/s, and the ratio P of the capillary viscosity (CV_D) of the carboxylic acid-modified nitrile-based copolymer latex in the de-swollen state to the capillary viscosity (CV_0) of the carboxylic acid-modified nitrile-based copolymer latex in the swollen state in the methyl ethyl ketone solvent is adjusted to a range of 0.85 to 1.0, such that elasticity and softness of a molded article produced by using the carboxylic acid-modified nitrile-based copolymer latex may be improved.

[Description of Drawings]

[0013]

FIG. 1 is a graph showing a change in pH according to an addition amount of KOH.

FIG. 2 is a derivative curve of the addition amount of KOH according to the pH.

[Best Mode]

[0014] The terms and words used in the description and claims of the present invention are not to be construed limitedly as having general or dictionary meanings but are to be construed as having meanings and concepts meeting the technical ideas of the present invention, based on a principle that the inventors are able to appropriately define the concepts of terms in order to describe their own inventions in the best mode.

[0015] The term "a monomer-derived unit" in the present invention may refer to a component, a structure, or a material itself derived from a monomer, and as a specific example, may refer to a unit formed in the polymer by the introduced monomer participating in the polymerization reaction during polymerization.

[0016] The term "latex" in the present invention may refer to a polymer or copolymer polymerized by polymerization being present in a form dispersed in water, and as a specific example, may refer to fine particles of a polymer in a rubber phase or a copolymer in a rubber phase polymerized by emulsion polymerization being present in a form dispersed in water in a colloid state.

[0017] The term "derived layer" in the present invention may represent a layer formed from a polymer or a copolymer, and as a specific example, a layer formed from a polymer or a copolymer by attaching, fixing, and/or polymerizing the polymer or the copolymer on a dip mold during manufacture of molded article by dip molding.

[0018] The term, "cross-linking agent-derived cross-linking part" in the present invention may refer to a component, a

structure, or a material itself caused by a compound, or a cross-linking part serving as a cross-linker in a polymer or between polymers, formed by the action or reaction of the cross-linking agent composition.

[0019] The term "alkyl" in the present invention may refer to a linear or branched saturated monovalent hydrocarbon of carbon atoms, such as, methyl, ethyl, propyl, 2-propyl, n-butyl, iso-butyl, tert-butyl, pentyl, hexyl, and dodecyl, or may include alkyl substituted by a substituent as well as unsubstituted alkyl.

[0020] The term "aryl" in the present invention may refer to an alkyl group as defined above in which one or more hydrogen atoms are substituted by an aryl group, such as phenyl, naphthalenyl, and fluorenyl, or may include aryl substituted by a substituent as well as unsubstituted aryl.

[0021] The term "(meth)acrylate" in the present invention may refer to both acrylate and methacrylate.

[0022] Hereinafter, the present invention will be described in more detail in order to assist in the understanding of the present invention.

[0023] According to the present invention, there is provided a latex composition for dip molding comprising a carboxylic acid-modified nitrile-based copolymer latex capable of improving elasticity and softness of a molded article. A carboxylic acid-modified nitrile-based copolymer in the carboxylic acid-modified nitrile-based copolymer latex may include a conjugated diene-based monomer-derived unit, an ethylenic unsaturated nitrile-based monomer-derived unit, and an ethylenic unsaturated acid monomer-derived unit.

[0024] The conjugated diene-based monomer constituting the conjugated diene-based monomer-derived unit may be one or more selected from the group consisting of 1,3-butadiene, 2,3-dimethyl-1,3-butadiene, 2-ethyl-1,3-butadiene, 1,3-pentadiene, and isoprene. As a specific example, the conjugated diene-based monomer may be 1,3-butadiene or isoprene.

[0025] A content of the conjugated diene-based monomer-derived unit may be 64.5 wt% to 83.5 wt%, or 67.5 wt% to 79.5 wt%, with respect to a total content of the carboxylic acid-modified nitrile-based copolymer. Within this range, a molded article formed of the latex composition for dip molding comprising the carboxylic acid-modified nitrile-based copolymer may be excellent in wearing comfort due to its high elasticity and softness.

[0026] The ethylenic unsaturated nitrile-based monomer constituting the ethylenic unsaturated nitrile-based monomer-derived unit may be one or more selected from the group consisting of acrylonitrile, methacrylonitrile, fumaronitrile, α -chloronitrile, and α -cyanoethylacrylonitrile. As a specific example, the ethylenic unsaturated nitrile-based monomer may be acrylonitrile or methacrylonitrile, and as a more specific example, the ethylenic unsaturated nitrile-based monomer may be acrylonitrile.

[0027] A content of the ethylenic unsaturated nitrile-based monomer-derived unit may be 15 wt% to 30 wt%, or 18 wt% to 28 wt%, with respect to the total content of the carboxylic acid-modified nitrile-based copolymer. Within this range, the molded article formed of the latex composition for dip molding comprising the carboxylic acid-modified nitrile-based copolymer may be excellent in wearing comfort due to its high elasticity and softness.

[0028] The ethylenic unsaturated acid monomer constituting the ethylenic unsaturated acid monomer-derived unit may be an ethylenic unsaturated monomer having an acid group such as a carboxyl group, a sulfonic acid group, or an acid anhydride group. As a specific example, the ethylenic unsaturated acid monomer may include one or more selected from the group consisting of an ethylenic unsaturated acid monomer such as acrylic acid, methacrylic acid, itaconic acid, maleic acid, or fumaric acid; polycarboxylic acid anhydride such as maleic acid anhydride or citraconic acid anhydride; an ethylenic unsaturated sulfonic acid monomer such as styrenesulfonic acid; and an ethylenic unsaturated polycarboxylic acid partial ester monomer such as monobutyl fumarate, monobutyl maleate, or mono-2-hydroxypropyl maleate. As a more specific example, the ethylenic unsaturated acid monomer may be one or more selected from the group consisting of acrylic acid, methacrylic acid, itaconic acid, maleic acid, and fumaric acid. As a still more specific example, the ethylenic unsaturated acid monomer may be methacrylic acid. The ethylenic unsaturated acid monomer may be used in a form of a salt such as an alkali metal salt or an ammonium salt during polymerization.

[0029] A content of the ethylenic unsaturated acid monomer-derived unit may be 1.5 wt% to 5.5 wt%, or 2.5 wt% to 4.5 wt%, with respect to the total content of the carboxylic acid-modified nitrile-based copolymer. Within this range, the molded article formed of the latex composition for dip molding comprising the carboxylic acid-modified nitrile-based copolymer may be excellent in wearing comfort due to its high elasticity and softness.

[0030] The carboxylic acid-modified nitrile-based copolymer latex of the present invention may further selectively include an ethylenic unsaturated monomer-derived unit, in addition to the conjugated diene-based monomer-derived unit, the ethylenic unsaturated nitrile-based monomer-derived unit, and the ethylenic unsaturated acid monomer-derived unit.

[0031] The ethylenic unsaturated monomer constituting the ethylenic unsaturated monomer-derived unit may be one or more selected from the group consisting of a hydroxyalkyl (meth)acrylate monomer having 1 to 4 carbon atoms; a vinyl aromatic monomer selected from the group consisting of styrene, alkylstyrene, and vinyl naphthalene; a fluoroalkyl vinyl ether monomer such as fluoro ethyl vinyl ether; an ethylenic unsaturated amide monomer selected from the group consisting of (meth)acrylamide, N-methylol (meth)acrylamide, N,N-dimethylol (meth)acrylamide, N-methoxymethyl (meth)acrylamide, and N-propoxymethyl (meth)acrylamide; a non-conjugated diene monomer such as vinyl pyridine,

vinyl norbornene, dicyclopentadiene, or 1,4-hexadiene; and an ethylenic unsaturated carboxylic acid ester monomer selected from the group consisting of methyl (meth)acrylate, ethyl (meth)acrylate, butyl (meth)acrylate, 2-ethylhexyl (meth)acrylate, trifluoroethyl (meth)acrylate, tetrafluoropropyl (meth)acrylate, dibutyl maleate, dibutyl fumarate, diethyl maleate, methoxymethyl (meth)acrylate, ethoxyethyl (meth)acrylate, methoxyethoxyethyl (meth)acrylate, cyanomethyl (meth)acrylate, 2-cyanoethyl (meth)acrylate, 1-cyanopropyl (meth)acrylate, 2-ethyl-6-cyanoethyl (meth)acrylate, 3-cyanopropyl (meth)acrylate, hydroxyethyl (meth)acrylate, hydroxypropyl (meth)acrylate, glycidyl (meth)acrylate, and dimethylamino ethyl (meth)acrylate.

[0032] In a case where the ethylenic unsaturated monomer-derived unit is selectively included in the carboxylic acid-modified nitrile-based copolymer, a content of the ethylenic unsaturated monomer-derived unit may be 0.5 wt% to 5 wt% with respect to the total content of the carboxylic acid-modified nitrile-based copolymer. Within this range, it is possible to improve properties such as a tensile strength of the molded article formed of the latex composition for dip molding comprising the carboxylic acid-modified nitrile-based copolymer.

[0033] The carboxylic acid-modified nitrile-based copolymer latex of the present invention has a monomer composition in which the contents of the ethylenic unsaturated nitrile-based monomer and the ethylenic unsaturated acid monomer are optimized, such that the elasticity and softness of the molded article formed of the carboxylic acid-modified nitrile-based copolymer latex are improved.

[0034] In the present invention, features of the latex that have never been applied before may be applied by controlling not only the monomer composition in the carboxylic acid-modified nitrile-based copolymer latex but also process factors of a preparation method described below.

[0035] Specifically, first, a preparation method of the carboxylic acid-modified nitrile-based copolymer latex includes a polymerization step of adding and polymerizing a conjugated diene-based monomer, an ethylenic unsaturated nitrile-based monomer, and an ethylenic unsaturated acid monomer into a polymerization reactor.

[0036] The polymerization step is performed to form a main chain of the carboxylic acid-modified nitrile-based copolymer latex, and the polymerization is performed by emulsion polymerization. In this case, the monomers are first added to the polymerization reactor prior to a polymerization reaction in the types and contents described above. The monomers may be separately added, added at the same time, or continuously added. For example, in a case where the monomers are added at the same time, the conjugated diene-based monomer, the ethylenic unsaturated nitrile-based monomer, and the ethylenic unsaturated acid monomer may be added to the polymerization reactor at the same time. In addition, in a case where the monomers are separately added, the monomers may be separately added by primarily adding a part of the monomers to the polymerization reactor and secondarily adding the remaining monomers to the polymerization reactor. As a specific example, in the primary addition, the ethylenic unsaturated nitrile-based monomer and the ethylenic unsaturated acid monomer may be added, and in the secondary addition, the conjugated diene-based monomer may be added.

[0037] In addition, in a case where the monomers are added prior to the polymerization reaction, additives such as an emulsifier and a chain transfer agent and a medium may be added at the same time or may be separately added after the monomers are added and stirring is started. For example, in the case where the monomers are added at the same time prior to the polymerization reaction, the additives and the medium may be added after the addition of the monomers. In the case where the monomers are separately added prior to the polymerization reaction, the additives and the medium may be added between the primary addition and the secondary addition. In a case where the monomers are first added prior to the addition of the additives and the medium and then stirring is performed, the composition and amounts of the monomers dissolved in an aqueous phase at the beginning of the polymerization reaction are affected, which affects a molecular weight of the carboxylic acid-modified nitrile-based copolymer to be polymerized or an acid distribution of the copolymer in latex particles.

[0038] Thereafter, in the polymerization step, the polymerization reaction is started by adding a polymerization initiator, and the conjugated diene-based monomer, the ethylenic unsaturated nitrile-based monomer, and the ethylenic unsaturated acid monomer may be added first, second, and third during the polymerization reaction, that is, may be separately added in the n^{th} order depending on the types of the monomers.

[0039] As such, in the case where the monomers are separately added by controlling the timing of addition of the monomers during the polymerization reaction, when a monomer-derived unit is formed from each monomer in the carboxylic acid-modified nitrile-based copolymer, a distribution of the monomers according to a difference in reaction speed of each monomer may be uniform. Therefore, a balance between physical properties of the molded article produced by using the carboxylic acid-modified nitrile-based copolymer may be improved.

[0040] In addition, in the case where the monomers are separately added during the polymerization reaction, the additives such as the emulsifier and the chain transfer agent and the medium may be separately added accordingly. In this case, the distribution of the monomers according to the difference in reaction speed of each monomer may be easily controlled, and a balance between physical properties of the molded article produced by using the carboxylic acid-modified nitrile-based copolymer may be improved.

[0041] One or more selected from the group consisting of an anionic surfactant, a nonionic surfactant, a cationic

surfactant, and an amphoteric surfactant may be used as the emulsifier. As a specific example, the emulsifier may be one or more anionic surfactants selected from the group consisting of alkyl benzene sulfonate, aliphatic sulfonate, a higher alcohol sulfate ester salt, α -olefin sulfonate, and alkyl ether sulfate. In addition, the emulsifier is added in an amount of 2 parts by weight to 4 parts by weight with respect to a total content of 100 parts by weight of the monomers to be added in the polymerization step. In general, when the amount of the emulsifier is large, a particle size of the particle of the carboxylic acid-modified nitrile-based copolymer is reduced, resulting in excellent stability, whereas, in the production process of the molded article, it is difficult to remove the emulsifier and the amount of foam generated is increased, which makes it difficult to produce the molded article.

[0042] The chain transfer agent may include one or more selected from the group consisting of α -methylstyrene dimer; mercaptans such as t-dodecyl mercaptan, n-dodecyl mercaptan, and octyl mercaptan; halogenated hydrocarbon such as carbon tetrachloride, methylene chloride, or methylene bromide; and a sulfur-containing compound such as tetraethylthiuram disulfide, dipentamethylenethiuram disulfide, or diisopropyl xanthogen disulfide. As a specific example, the chain transfer agent may be mercaptans. The chain transfer agent may be added in an amount of 0.2 parts by weight to 0.9 parts by weight with respect to the total content of 100 parts by weight of the monomers to be added in the polymerization step. Within the above range, physical properties of the molded article may be excellent.

[0043] As a specific example, deionized water may be used as the medium. A temperature of the water used as the medium during the polymerization of the carboxylic acid-modified nitrile-based copolymer may be 45°C to 80°C. In general, emulsion polymerization is performed using water at room temperature as polywater. However, in a case where the high-temperature water is used as polywater, the composition and amounts of the monomers dissolved in the aqueous phase and the amounts of the emulsifier and the chain transfer agent are affected, which affects the acid distribution of the copolymer to be polymerized in the carboxylic acid-modified nitrile-based copolymer latex particles or an entanglement degree of the copolymer. In addition, the medium may be added in an amount of 105 parts by weight to 140 parts by weight with respect to the total content of 100 parts by weight of the monomers to be added in the polymerization step.

[0044] When the carboxylic acid-modified nitrile-based copolymer is polymerized, an additive such as a chelating agent, a dispersant, a pH adjuster, a deoxidizer, a particle size adjuster, an anti-aging agent, or an oxygen scavenger may be further added, if necessary.

[0045] The polymerization reaction may be started by adding a polymerization initiator.

[0046] A radical initiator may be used as the polymerization initiator. It is possible to use one or more selected from the group consisting of inorganic peroxide such as sodium persulfate, potassium persulfate, ammonium persulfate, potassium perphosphate, or hydrogen peroxide; organic peroxide such as diisopropylbenzene hydroperoxide, t-butyl hydroperoxide, cumene hydroperoxide, p-menthane hydroperoxide, di-t-butyl peroxide, t-butylcumyl peroxide, acetyl peroxide, isobutyl peroxide, octanoyl peroxide, benzoyl peroxide, 3,5,5-trimethylhexanol peroxide, or t-butyl peroxy isobutylate; and a nitrogen compound such as azobisisobutyronitrile, azobis-2,4-dimethylvaleronitrile, azobiscyclohexanecarbonitrile, or methyl azobis(isobutyrate). As a specific example, the polymerization initiator may be inorganic peroxide, and as a more specific example, the polymerization initiator may be a persulfate initiator. In addition, the polymerization initiator may be added in an amount of 0.1 parts by weight to 0.5 parts by weight with respect to the total content of 100 parts by weight of the monomers to be added in the polymerization step. Within this range, a polymerization speed may be maintained at an appropriate level.

[0047] Here, in a case where the organic peroxide or the inorganic peroxide is used as the polymerization initiator, the organic peroxide or the inorganic peroxide may be used in combination with an activator. As the activator, a compound containing metal ions in a reduced state, such as ferrous sulfate and cuprous naphthenate; a sulfonic acid compound such as sodium methanesulfonate; and an amine compound such as dimethylaniline may be used alone or in combination of two or more thereof. Specifically, as the activator, it is possible to use one or more selected from the group consisting of sodium formaldehyde, sulfoxylate, sodium ethylenediaminetetraacetate, ferrous sulfate, dextrose, sodium pyrophosphate, and sodium sulfite. The activator may be added in an amount of 0.05 parts by weight to 1 part by weight with respect to the total content of 100 parts by weight of the monomers to be added in the polymerization step. Within this range, the polymerization speed may be maintained at an appropriate level.

[0048] The polymerization of the carboxylic acid-modified nitrile-based copolymer may be performed at a temperature of 5°C to 60°C. Within this range, the polymerization speed may be maintained at an appropriate level.

[0049] In addition, the polymerization of the carboxylic acid-modified nitrile-based copolymer may be performed by controlling a change in reaction pressure relative to an initial reaction pressure according to a polymerization conversion rate to be in a specific range. Specifically, the reaction pressure at the beginning of the polymerization may be in a range of 2.0 kgf/cm² to 2.8 kgf/cm², and the reaction pressure when the polymerization conversion rate is 1% to 45% may be controlled in a range of 15% or less of an increase amount of the pressure relative to the reaction pressure at the beginning of the polymerization. In particular, the reaction pressure when the polymerization conversion rate is 40% may be in a range of 5% to 10% of the increase amount of the pressure relative to the reaction pressure at the beginning of the polymerization.

[0050] The reaction pressure when the polymerization conversion rate is 46% to 75% may be controlled in a range

of 5% to 70% of the increase amount of the pressure relative to the reaction pressure at the beginning of the polymerization. In particular, the reaction pressure when the polymerization conversion rate is 60% may be in a range of 30% to 65% of the increase amount of the pressure relative to the reaction pressure at the beginning of the polymerization.

[0051] The reaction pressure from the time when the polymerization conversion rate is 76% until the polymerization is terminated may be controlled in a range of 0% to 5% of the increase amount of the pressure relative to the reaction pressure at the beginning of the polymerization or in a range of 0% to 100% of a decrease amount of the pressure relative to the reaction pressure at the beginning of the polymerization. In particular, the reaction pressure when the polymerization conversion rate is 90% may be adjusted to a range of 10% or more of the decrease amount of the pressure relative to the reaction pressure at the beginning of the polymerization.

[0052] In the polymerization reaction of the carboxylic acid-modified nitrile-based copolymer, the polymerization conversion rate may be measured by a method commonly known in the art. For example, a predetermined amount of sample is collected from a reaction composition at a predetermined time interval, a solid content is measured, and then, a polymerization conversion rate is calculated according to the following Equation 1.

[Equation 1]

$$\begin{aligned} \text{Polymerization conversion rate (\%)} = & \left[\left\{ \left(\text{parts by weight of added monomers and additives} \right) \times \text{solid content (\%)} \right. \right. \\ & \left. \left. - \left(\text{parts by weight of additives added other than monomers} \right) \right\} / \left(\text{total parts by weight of added monomers} \right) \right] \times 100 \end{aligned}$$

[0053] In Equation 1, parts by weight are based on the total content of 100 parts by weight of the monomers to be added. A sectional polymerization conversion rate may be calculated by reflecting parts by weight of the monomers and additives added until a corresponding section.

[0054] When the change in reaction pressure relative to the initial reaction pressure according to the polymerization conversion rate is adjusted to be in the above range, the composition and amounts of the monomers dissolved in the aqueous phase during the polymerization reaction are affected depending on the reaction pressure, which affects an entanglement degree or a branch degree of the carboxylic acid-modified nitrile-based copolymer to be polymerized.

[0055] The preparation method of the carboxylic acid-modified nitrile-based copolymer latex includes a step of terminating the polymerization reaction and obtaining a carboxylic acid-modified nitrile-based copolymer latex. The polymerization reaction is terminated by cooling the polymerization system at the time when the polymerization conversion rate is 85% or more, or adding a polymerization terminator, a pH adjuster, or an antioxidant.

[0056] In addition, the preparation method of the carboxylic acid-modified nitrile-based copolymer latex may include a step of removing unreacted monomers by a deodorization process after the reaction is terminated.

[0057] In the present invention, physical properties of the latex itself may be improved to improve physical properties such as softness and elasticity of a molded article produced by a dip molding process using a latex composition for dip molding comprising a carboxylic acid-modified nitrile-based copolymer latex. Specifically, in the composition of the monomers in the latex, the amounts of the ethylenic unsaturated nitrile-based monomer and the ethylenic unsaturated acid monomer are optimized, and in the polymerization step, the entanglement degree or the branch degree and the molecular weight of the carboxylic acid-modified nitrile-based copolymer and the acid distribution of the copolymer in the latex particles are adjusted, thereby optimizing a structure of the latex particles.

[0058] Such a carboxylic acid-modified nitrile-based copolymer may be prepared by the composition and polymerization method of the copolymer described above. In the present invention, a numerical range of each parameter that satisfies the physical properties required by the present inventors is determined by using the following various parameters while adjusting the physical properties of the carboxylic acid-modified nitrile-based copolymer.

[0059] According to an exemplary embodiment of the present invention, the carboxylic acid-modified nitrile-based copolymer latex may satisfy the following General Formula 1 and General Formula 2.

[General Formula 1]

$$1.0 \text{ mm}^2/\text{s} \leq CV_0 \leq 3.0 \text{ mm}^2/\text{s}$$

[General Formula 2]

$$0.85 \leq P \leq 1.0, P = CV_D / CV_0$$

[0060] In General Formulas 1 and 2, CV_0 represents a capillary viscosity of the carboxylic acid-modified nitrile-based copolymer latex measured in a swollen state in a methyl ethyl ketone solvent, and CV_D represents a capillary viscosity of the carboxylic acid-modified nitrile-based copolymer latex in a de-swollen state.

[0061] In General Formula 1, CV_0 is measured using a Cannon-Fenske routine type capillary viscometer (SI Analytics GmbH Type No. 520 13) and a methyl ethyl ketone (MEK) solvent under a condition of a pH of 8.2 to 9.2. Accordingly, CV_0 represents a capillary viscosity of the carboxylic acid-modified nitrile-based copolymer latex measured in a swollen state in the methyl ethyl ketone solvent. In general, the capillary viscosity is used as a means for measuring a molecular weight of a non-cross-linked polymer. However, in the present invention, CV_0 represents a capillary viscosity measured in a state where the copolymer particles in the latex are swollen in the methyl ethyl ketone solvent, that is, a swollen state of the copolymer particles in the latex, unlike such a general capillary viscosity. Therefore, it is possible to acquire, from CV_0 , information on not only the molecular weight of the copolymer in the carboxylic acid-modified nitrile-based copolymer latex but also factors affecting the swelling by the methyl ethyl ketone solvent, and among this information, in particular, information on the acid distribution of the copolymer in the latex particles may be efficiently acquired. CV_0 may be 0.5 to 4 or 1 to 3.

[0062] In order to improve softness of the molded article produced by using the latex composition for dip molding required in the present invention, it is required for the carboxylic acid-modified nitrile-based copolymer to be as soft and stretchable as possible. To this end, the molecular weight of the copolymer is required to be optimized. When the molecular weight of the copolymer is not optimized, the copolymer becomes stiff and hard or the copolymer does not stretch well. In addition, when preparing the latex composition for dip molding, the ethylenic unsaturated acid monomer-derived unit in the carboxylic acid-modified nitrile-based copolymer latex may form an ion bond with metal ions to be added in a metal oxide form. Such a degree of bonding with the metal ions may also have a great influence on the softness of the molded article produced by using the latex composition for dip molding. The degree of bonding with the metal ions has an important influence on the structure of the latex particles, such as the acid distribution of the copolymer in the latex particles. Therefore, when a molded article is produced by using the carboxylic acid-modified nitrile-based copolymer having the optimal acid distribution of the copolymer in the latex particles, softness of the molded article may be improved.

[0063] In this regard, since the carboxylic acid-modified nitrile-based copolymer latex of the present invention satisfying the range of CV_0 has an appropriate level of the molecular weight of the copolymer and the optimal acid distribution of the copolymer in the latex particles, when the carboxylic acid-modified nitrile-based copolymer latex is used in the latex composition for dip molding, it is possible to implement the molded article having excellent physical properties, particularly, excellent softness.

[0064] In General Formula 2, CV_0 represents a capillary viscosity of the carboxylic acid-modified nitrile-based copolymer latex measured in a swollen state in a methyl ethyl ketone solvent, and CV_D represents a capillary viscosity of the carboxylic acid-modified nitrile-based copolymer latex in a de-swollen state, as described in General Formula 1.

[0065] Specifically, a state where the copolymer particles in the latex are swollen in the methyl ethyl ketone solvent may be defined as the swollen state, and the de-swollen state may refer to a state where a swollen portion of the copolymer particles in the latex in the methyl ethyl ketone solvent is removed. For example, a state where a portion of the copolymer particles dissolved in the methyl ethyl ketone solvent is removed by applying a predetermined energy to the copolymer particles in the latex in the swollen state may refer to the de-swollen state. CV_D may be a capillary viscosity measured by dissolving the carboxylic acid-modified nitrile-based copolymer latex in the methyl ethyl ketone solvent under a condition of a pH of 8.2 to 9.2, and de-swelling the carboxylic acid-modified nitrile-based copolymer latex by applying an energy of 55 kcal to 65 kcal using an ultrasonicator (Branson[®] M Mechanical Bath 5800) for 40 minutes. In the case, the capillary viscosity may be measured in the same method as that in the measurement of CV_0 using the Cannon-Fenske routine type capillary viscometer (SI Analytics GmbH Type No. 520 13).

[0066] In addition, P may refer to a ratio of CV_0 and CV_D . Specifically, P may refer to a ratio of CV_D to CV_0 . More specifically, P may refer to a ratio of the capillary viscosity of the carboxylic acid-modified nitrile-based copolymer latex in the de-swollen state to the capillary viscosity of the carboxylic acid-modified nitrile-based copolymer latex in the swollen state in the methyl ethyl ketone solvent. P may be 0.85 to 1 or 0.9 to 1. When a value of P of the carboxylic acid-modified nitrile-based copolymer latex is within the above range, it may mean that a change in capillary viscosity is small even when the carboxylic acid-modified nitrile-based copolymer latex is de-swollen by applying an additional energy. This may mean that the carboxylic acid-modified nitrile-based copolymer latex has a structure in which the portion dissolved in the methyl ethyl ketone solvent is relatively small and the portion to be dissolved when being de-swollen is difficult to escape. That is, this may mean the entanglement degree of the copolymer particles in the latex is relatively large and

the carboxylic acid-modified nitrile-based copolymer latex has a latex particle structure in which an entangled copolymer is hardly untangled. Accordingly, it is possible to acquire, from P, information on the latex particle structure such as the entanglement degree or the branch degree of the copolymer in the carboxylic acid-modified nitrile-based copolymer latex.

[0067] In order to improve elasticity, that is, a stress retention rate of the molded article produced by using the latex composition for dip molding required in the present invention, there is a need for a carboxylic acid-modified nitrile-based copolymer latex having a structure in which the entanglement degree or branch degree of the copolymer in the carboxylic acid-modified nitrile-based copolymer latex is increased. This is for the purpose of imparting the original elasticity of rubber to the latex, and, for example, even in a case where cross-linking is performed with a cross-linking agent that remarkably increases the stress retention rate, a stress retention rate of the existing latex having a linear chain is not increased because it does not inherently have elasticity. On the other hand, since the latex of the present invention having a structure in which the entanglement degree or the branch degree is increased has rubber elasticity, when this latex is used in a latex composition for dip molding, elasticity of a molded article, that is, a stress retention rate may be improved.

[0068] Therefore, the carboxylic acid-modified nitrile-based copolymer latex of the present invention satisfying the range of P has an appropriate level of the entanglement degree of the copolymer and a latex particle structure in which the entangled copolymer is hardly untangled. When this carboxylic acid-modified nitrile-based copolymer latex is used in a latex composition for dip molding, elasticity of a molded article, that is, a stress retention rate may be improved.

[0069] According to an exemplary embodiment of the present invention, the carboxylic acid-modified nitrile-based copolymer latex may satisfy the following General Formula 3 and General Formula 4.

[General Formula 3]

$$5 \leq M \leq 20, M = m_1 \times m_2$$

[General Formula 4]

$$9.2 \leq \text{pKa} \leq 10.5$$

[0070] In General Formula 3, m_1 represents an insolubility of a dried film of the carboxylic acid-modified nitrile-based copolymer latex in the methyl ethyl ketone solvent, and m_2 represents a swelling index of the dried film of the carboxylic acid-modified nitrile-based copolymer latex in the methyl ethyl ketone solvent.

[0071] In addition, in General Formula 3, a value of M is a value expressed as a product of the insolubility of the dried film of the carboxylic acid-modified nitrile-based copolymer latex in the methyl ethyl ketone solvent and the swelling index of the dried film in the methyl ethyl ketone solvent, and the value may satisfy a range of 5.0 to 20 or 7 to 15. The insolubility of the dried film of the carboxylic acid-modified nitrile-based copolymer latex in the methyl ethyl ketone solvent is an index indicating a degree of cross-linking of the carboxylic acid-modified nitrile-based copolymer contained in the carboxylic acid-modified nitrile-based copolymer latex. The swelling index of the dried film of the carboxylic acid-modified nitrile-based copolymer latex in the methyl ethyl ketone solvent is an index indicating swellability of the carboxylic acid-modified nitrile-based copolymer contained in the carboxylic acid-modified nitrile-based copolymer latex in the methyl ethyl ketone solvent. Therefore, from M obtained by multiplying these two indices, it is possible to acquire information on the entanglement degree of the carboxylic acid-modified nitrile-based copolymer latex or the acid distribution of the copolymer in the latex particles.

[0072] In General Formula 4, a value of pKa is an index indicating a degree of ionization of an ethylenic unsaturated acid of the carboxylic acid-modified nitrile-based copolymer latex, and information on the acid distribution of the copolymer in the latex particles may be acquired from the value of pKa. The value of pKa may be in a range of 9.2 to 10.5 or 9.5 to 10.2.

[0073] The carboxylic acid-modified nitrile-based copolymer latex satisfying the ranges of M and pKa has an appropriate level of the entanglement degree of the copolymer and the optimal acid distribution of the copolymer in the latex particles.

Therefore, in a case where the carboxylic acid-modified nitrile-based copolymer latex is used in a latex composition for dip molding, in the production process of the molded article, an occurrence of a flow mark is prevented, such that workability may be improved, and polymerization stability during polymerization of the carboxylic acid-modified nitrile-based copolymer latex may be improved.

[0074] As such, the latex is prepared while confirming the numerical ranges of CVo, P, M, and pKa, thereby completing the latex of the present invention.

[0075] According to an exemplary embodiment of the present invention, the latex composition for dip molding may comprise a carboxylic acid-modified nitrile-based copolymer latex and a cross-linking agent composition. The cross-linking agent composition may be used to form a cross-linking agent-derived cross-linking part through a cross-linking

reaction of the carboxylic acid-modified nitrile-based copolymer.

[0076] The cross-linking agent composition may include one or more selected from the group consisting of a vulcanizing agent, a vulcanization accelerator, a divalent cationic metal oxide, and a polyvalent metal cationic compound.

[0077] The vulcanizing agent is used for vulcanizing the latex composition for dip molding, and may be sulfur, and as a specific example, the vulcanizing agent may be sulfur such as powdered sulfur, precipitated sulfur, colloidal sulfur, surface-treated sulfur, or insoluble sulfur. A content of the vulcanizing agent may be 0 parts by weight to 3 parts by weight with respect to the total content of 100 parts by weight of the carboxylic acid-modified nitrile-based copolymer in the latex composition for dip molding. Within this range, cross-linking ability by vulcanization may be excellent.

[0078] In addition, the vulcanization accelerator may be one or more selected from the group consisting of 2-mercaptobenzothiazole (MBT), 2,2-dithiobisbenzothiazole-2-sulfenamide (MBTS), N-cyclohexylbenzothiazole-2-sulfenamide (CBS), 2-morpholinothiobenzothiazole (MBS), tetramethylthiuram monosulfide (TMTM), tetramethylthiuram disulfide (TMTD), zinc diethyldithiocarbamate (ZDEC), zinc di-n-butylthiocarbamate (ZDBC), diphenylguanidine (DPG), and di-o-tolylguanidine. A content of the vulcanization accelerator may be 0 parts by weight to 3 parts by weight with respect to the total content of 100 parts by weight of the carboxylic acid-modified nitrile-based copolymer in the latex composition for dip molding. Within this range, cross-linking ability by vulcanization may be excellent.

[0079] In addition, the divalent cationic metal oxide may include one or more selected from the group consisting of divalent cationic metals, for example, oxides such as Mg, Ca, Zn, Sr, and Ba. As a preferred example, the divalent cationic metal oxide may include one or more selected from the group consisting of zinc oxide, magnesium oxide, and calcium oxide.

[0080] The divalent cationic metal oxide may be a cross-linking agent for forming a cross-linking part through ionic bonding in the carboxylic acid-modified nitrile-based copolymer or between carboxylic acid-modified nitrile-based copolymers by performing ionic bonding with a functional group of the ethylenic unsaturated acid monomer of the carboxylic acid-modified nitrile-based copolymer in the latex composition for dip molding. A content of the divalent cationic metal oxide may be 0.1 parts by weight to 3 parts by weight with respect to the total content of 100 parts by weight of the carboxylic acid-modified nitrile-based copolymer in the latex composition for dip molding. Within this range, ionic bonding ability is excellent and a tensile strength of the produced dip-molded article may be excellent.

[0081] In addition, the polyvalent metal cationic compound may include one or more selected from the group consisting of a polyvalent metal cation having a charge of +2 or higher and a ligand.

[0082] The polyvalent metal cation may include, for example, one or more selected from the group consisting of a polyvalent cation of an alkaline earth metal and a polyvalent cation of a transition metal.

[0083] The ligand is used to allow the polyvalent metal cation to be slowly eluted, and may include one or more selected from the group consisting of acetylacetonate, acetylacetate, glycolate, citrate, tartrate, gluconate, and nitriloacetate.

[0084] As a specific example, the polyvalent metal cationic compound may be an aluminum compound containing a complex anion such as acetylacetonate, acetylacetate, glycolate, citrate, tartrate, gluconate, or nitriloacetate.

[0085] A use amount of the ligand may be 1 equivalent to 3 equivalents of a use amount of the polyvalent metal cation. For example, in a case where the aluminum compound containing a complex anion is used as the polyvalent metal cation, the use amount of the ligand that allows the aluminum ions to be slowly eluted may be 1 equivalent to 3 equivalents of a use amount of aluminum. As described above, in a case where the polyvalent metal cationic compound is used as a cross-linking agent for ionic bonding or covalent bonding, it is possible to prevent bonding in an acidic solution from being weakened, which may prevent a deterioration of the strength of the dip-molded article. In addition, by using the cross-linking agent composition containing the polyvalent metal cationic compound, cross-linking is formed through ionic bonding with the acids distributed in the latex particles, such that it is possible to prevent a deterioration of the physical properties of the molded article caused by the absence of sulfur and a vulcanization accelerator. In addition, in this case, the dip-molded article may be produced without addition of a vulcanization accelerator. Thus, it is possible to prevent side effects caused to users due to the vulcanization accelerator.

[0086] A content of the polyvalent metal cationic compound may be 0.1 parts by weight to 3 parts by weight with respect to 100 parts by weight (based on a solid content) of the carboxylic acid-modified nitrile-based copolymer latex composition. Within this range, it is possible to prevent a deterioration of the strength of the dip-molded article.

[0087] In addition, according to an exemplary embodiment of the present invention, the latex composition for dip molding may further include an additive such as a dispersant, an antioxidant, a pigment, a filler, a thickening agent, or a pH adjuster, if necessary.

[0088] According to an exemplary embodiment of the present invention, a solid content (concentration) of the latex composition for dip molding may be, for example, 5 wt% to 43 wt%. Within this range, latex transport efficiency may be excellent and an increase in viscosity of the latex may be prevented, such that storage stability may be excellent.

[0089] As another example, a pH of the latex composition for dip molding may be 9 to 11. Within this range, processability and productivity when producing a dip-molded article may be excellent. The pH of the latex composition for dip molding may be adjusted by addition of the pH adjuster described above. Examples of the pH adjuster may include an aqueous potassium hydroxide solution having a concentration of 1 wt% to 10 wt%, and ammonia water having a concentration

of 1 wt% to 50 wt%.

[0090] According to the present invention, a molded article including a latex composition for dip molding-derived layer is provided. The molded article may be a dip-molded article produced by dip-molding the latex composition for dip molding.

[0091] A method of producing the molded article may include a step of dipping the latex composition for dip molding by a direct dipping method, an anode coagulation dipping method, a Teague's coagulation dipping method, or the like. As a specific example, the molded article may be produced by an anode coagulation dipping method, and in this case, a dip-molded article having a uniform thickness may be obtained.

[0092] As a specific example, the method of producing the molded article may include: a step S100 of attaching a coagulant to a dip molding mold; a step S200 of dipping the dip molding mold to which the coagulant is attached into a latex composition for dip molding to form the latex composition for dip molding-derived layer, that is, a dip-molded layer; and a step S300 of heating the dip-molded layer to cross-link the latex composition for dip molding.

[0093] The step S100 is a step of dipping a dip molding mold into a coagulant solution to attach a coagulant onto a surface of the dip molding mold so that the coagulant is formed on the dip molding mold. The coagulant solution is a solution obtained by dissolving a coagulant in water, alcohol, or a mixture thereof. A content of the coagulant in the coagulant solution may be 5 wt% to 45 wt% with respect to a total content of the coagulant solution.

[0094] The coagulant may be, for example, one or more selected from the group consisting of metal halide such as barium chloride, calcium chloride, magnesium chloride, zinc chloride, or aluminum chloride; nitrate such as barium nitrate, calcium nitrate, or zinc nitrate; acetate such as barium acetate, calcium acetate, or zinc acetate; and sulfate such as calcium sulfate, magnesium sulfate, or aluminum sulfate. As a specific example, the coagulant may be calcium chloride or calcium nitrate.

[0095] In addition, the step S200 may be a step of dipping the dip molding mold to which the coagulant is attached into the latex composition for dip molding according to the present invention and taking out the dip molding mold to form a dip-molded layer on the dip molding mold.

[0096] In addition, the step S300 may be a step of heating the dip-molded layer formed on the dip molding mold and cross-linking and curing the latex composition for dip molding to obtain a dip-molded article.

[0097] Thereafter, the dip-molded layer cross-linked by the heating treatment may be removed from the dip molding mold to obtain a dip-molded article.

[0098] A modulus at 500% of the molded article may be 4 to 20.5 or 5 to 18.5. The modulus at 500% is a value indicating how much force is required to stretch the molded article to 500% of its original length. When the molded article has the value of the modulus at 500% in the above range, the molded article may be excellent in softness.

[0099] The molded article may have a stress retention rate of 40% to 70% or 40% to 65%. The stress retention rate is a value of a stress when an elongation of 100% is maintained for 5 minutes to a stress when an elongation is increased to 100%, expressed as a percentage, and within the range of the stress retention rate, the molded article may have excellent elasticity.

[0100] According to an exemplary embodiment of the present invention, the molded article may be a glove such as a surgical glove, an examination glove, an industrial glove, or a household glove, a condom, a catheter, or a health care product.

[0101] Hereinafter, the present invention will be described in more detail by examples. However, the following examples are provided for illustrating the present invention. It is apparent to a person skilled in the art that various modifications and alterations may be made without departing from the scope and spirit of the present invention, and the scope of the present invention is not limited thereto.

Examples

Example 1

<Preparation of carboxylic acid-modified nitrile-based copolymer latex>

[0102] Hereinafter, wt% indicates a total content of monomers to be added to form a main chain of a carboxylic acid-modified nitrile-based copolymer, and part(s) of weight indicates a total of 100 parts by weight of the monomers to be added.

[0103] 23 wt% of acrylonitrile and 3.5 wt% of methacrylic acid were added to a polymerization reactor installed with a stirrer and stirring was started to mix them, 0.6 parts by weight of t-dodecylmercaptan, 4.0 parts by weight of sodium dodecylbenzene sulfonate, and 120 parts by weight of water at 70°C were added, 73.5 wt% of 1,3-butadiene was added, and then, emulsion polymerization was initiated by adding 0.3 parts by weight of potassium persulfate as a polymerization initiator at a pressure of 2.5 kgf/cm² and a temperature of 40°C.

[0104] The reaction pressure at the time when the polymerization conversion rate was 40% was controlled to 2.68 kgf/cm², the reaction pressure at the time when the polymerization conversion rate was 60% was controlled to 3.55

kgf/cm², and the reaction pressure at the time when the polymerization conversion rate was 90% was controlled to 1.25 kgf/cm².

[0105] When the polymerization conversion rate reached 94%, the polymerization was terminated. Thereafter, unreacted products were removed through a deodorization process, and ammonium water, an antioxidant, and a defoaming agent were added, thereby obtaining a carboxylic acid-modified nitrile-based copolymer latex.

<Preparation of latex composition for dip molding>

[0106] To 100 parts by weight (based on a solid content) of the obtained carboxylic acid-modified nitrile-based copolymer latex, 1 part by weight of sulfur, 1.0 part by weight of zinc di-n-butylthiocarbamate (ZDBC), 1.3 parts by weight of zinc oxide, 1 part by weight of titanium oxide, an aqueous potassium hydroxide solution, and secondary distilled water were added to obtain a latex composition for dip molding having a solid content concentration of 30 wt% and a pH of 10.2.

<Production of molded article>

[0107] 18 wt% of calcium nitrate, 81.5 wt% of water, and 0.5 wt% of a wetting agent (Teric 320, produced by Huntsman Corporation, Australia) were mixed with each other to prepare a coagulant solution. A hand-shaped ceramic mold was dipped into the solution for 13 seconds and taken out from the solution, and the hand-shaped ceramic mold was dried at 120°C for 3 minutes, thereby applying a coagulant to the hand-shaped mold.

[0108] Subsequently, the hand-shaped mold to which the coagulant was applied was dipped into the obtained latex composition for dip molding for 13 seconds and taken out from the latex composition, the hand-shaped mold was dried at 80°C for 1 minute, and the dried hand-shaped mold was dipped into water or warm water for 60 seconds. Thereafter, cross-linking was performed at 120°C for 20 minutes. A cross-linked dip-molded layer was removed from the hand-shaped mold, thereby obtaining a molded article having a glove shape.

Example 2

[0109] 18 wt% of acrylonitrile and 4.5 wt% of methacrylic acid were added to a polymerization reactor installed with a stirrer and stirring was started to mix them, 0.3 parts by weight of t-dodecylmercaptan, 3.0 parts by weight of sodium dodecylbenzene sulfonate, and 130 parts by weight of water at 70°C were added, 77.5 wt% of 1,3-butadiene was added, and then, emulsion polymerization was initiated by adding 0.25 parts by weight of potassium persulfate as a polymerization initiator at a pressure of 2.7 kgf/cm² and a temperature of 40°C.

[0110] The reaction pressure at the time when the polymerization conversion rate was 40% was controlled to 2.92 kgf/cm², the reaction pressure at the time when the polymerization conversion rate was 60% was controlled to 4.27 kgf/cm², and the reaction pressure at the time when the polymerization conversion rate was 90% was controlled to 1.76 kgf/cm².

[0111] When the polymerization conversion rate reached 96%, the polymerization was terminated. Thereafter, unreacted products were removed through a deodorization process, and ammonium water, an antioxidant, and a defoaming agent were added, thereby obtaining a carboxylic acid-modified nitrile-based copolymer latex.

[0112] A latex composition for dip molding and a molded article were produced by using the obtained carboxylic acid-modified nitrile-based copolymer latex in the same manner as that of Example 1.

Example 3

[0113] 28 wt% of acrylonitrile and 2.5 wt% of methacrylic acid were added to a polymerization reactor installed with a stirrer and stirring was started to mix them, 0.8 parts by weight of t-dodecylmercaptan, 3.0 parts by weight of sodium dodecylbenzene sulfonate, and 110 parts by weight of water at 50°C were added, 69.5 wt% of 1,3-butadiene was added, and then, emulsion polymerization was initiated by adding 0.35 parts by weight of potassium persulfate as a polymerization initiator at a pressure of 2.2 kgf/cm² and a temperature of 37°C.

[0114] The reaction pressure at the time when the polymerization conversion rate was 40% was controlled to 2.38 kgf/cm², the reaction pressure at the time when the polymerization conversion rate was 60% was controlled to 3.17 kgf/cm², and the reaction pressure at the time when the polymerization conversion rate was 90% was controlled to 1.76 kgf/cm².

[0115] When the polymerization conversion rate reached 94%, the polymerization was terminated. Thereafter, unreacted products were removed through a deodorization process, and ammonium water, an antioxidant, and a defoaming agent were added, thereby obtaining a carboxylic acid-modified nitrile-based copolymer latex.

[0116] A latex composition for dip molding and a molded article were produced by using the obtained carboxylic acid-modified nitrile-based copolymer latex in the same manner as that of Example 1.

Example 4

[0117] 25 wt% of acrylonitrile and 4.1 wt% of methacrylic acid were added to a polymerization reactor installed with a stirrer and stirring was started to mix them, 0.3 parts by weight of t-dodecylmercaptan, 2.5 parts by weight of sodium dodecylbenzene sulfonate, and 140 parts by weight of water at 55°C were added, 70.9 wt% of 1,3-butadiene was added, and then, emulsion polymerization was initiated by adding 0.2 parts by weight of potassium persulfate as a polymerization initiator at a pressure of 2.4 kgf/cm² and a temperature of 40°C.

[0118] The reaction pressure at the time when the polymerization conversion rate was 40% was controlled to 2.52 kgf/cm², the reaction pressure at the time when the polymerization conversion rate was 60% was controlled to 3.12 kgf/cm², and the reaction pressure at the time when the polymerization conversion rate was 90% was controlled to 0.24 kgf/cm².

[0119] When the polymerization conversion rate reached 96%, the polymerization was terminated. Thereafter, unreacted products were removed through a deodorization process, and ammonium water, an antioxidant, and a defoaming agent were added, thereby obtaining a carboxylic acid-modified nitrile-based copolymer latex.

[0120] A latex composition for dip molding and a molded article were produced by using the obtained carboxylic acid-modified nitrile-based copolymer latex in the same manner as that of Example 1.

Example 5

[0121] 21 wt% of acrylonitrile and 2.9 wt% of methacrylic acid were added to a polymerization reactor installed with a stirrer and stirring was started to mix them, 0.6 parts by weight of t-dodecylmercaptan, 3.5 parts by weight of sodium dodecylbenzene sulfonate, and 110 parts by weight of water at 75°C were added, 76.1 wt% of 1,3-butadiene was added, and then, emulsion polymerization was initiated by adding 0.3 parts by weight of potassium persulfate as a polymerization initiator at a pressure of 2.6 kgf/cm² and a temperature of 39°C.

[0122] The reaction pressure at the time when the polymerization conversion rate was 40% was controlled to 2.86 kgf/cm², the reaction pressure at the time when the polymerization conversion rate was 60% was controlled to 4.29 kgf/cm², and the reaction pressure at the time when the polymerization conversion rate was 90% was controlled to 2.34 kgf/cm².

[0123] When the polymerization conversion rate reached 95%, the polymerization was terminated. Thereafter, unreacted products were removed through a deodorization process, and ammonium water, an antioxidant, and a defoaming agent were added, thereby obtaining a carboxylic acid-modified nitrile-based copolymer latex.

[0124] A latex composition for dip molding and a molded article were produced by using the obtained carboxylic acid-modified nitrile-based copolymer latex in the same manner as that of Example 1.

Example 6

[0125] 28 wt% of acrylonitrile and 4.5 wt% of methacrylic acid were added to a polymerization reactor installed with a stirrer and stirring was started to mix them, 0.25 parts by weight of t-dodecylmercaptan, 3.0 parts by weight of sodium dodecylbenzene sulfonate, and 110 parts by weight of water at 60°C were added, 67.5 wt% of 1,3-butadiene was added, and then, emulsion polymerization was initiated by adding 0.45 parts by weight of potassium persulfate as a polymerization initiator at a pressure of 2.0 kgf/cm² and a temperature of 36°C.

[0126] The reaction pressure at the time when the polymerization conversion rate was 40% was controlled to 2.2 kgf/cm², the reaction pressure at the time when the polymerization conversion rate was 60% was controlled to 2.6 kgf/cm², and the reaction pressure at the time when the polymerization conversion rate was 90% was controlled to 1.8 kgf/cm².

[0127] When the polymerization conversion rate reached 97%, the polymerization was terminated. Thereafter, unreacted products were removed through a deodorization process, and ammonium water, an antioxidant, and a defoaming agent were added, thereby obtaining a carboxylic acid-modified nitrile-based copolymer latex.

[0128] A latex composition for dip molding and a molded article were produced by using the obtained carboxylic acid-modified nitrile-based copolymer latex in the same manner as that of Example 1.

Example 7

[0129] 18 wt% of acrylonitrile and 2.5 wt% of methacrylic acid were added to a polymerization reactor installed with a stirrer and stirring was started to mix them, 0.5 parts by weight of t-dodecylmercaptan, 3.5 parts by weight of sodium dodecylbenzene sulfonate, and 120 parts by weight of water at 65°C were added, 79.5 wt% of 1,3-butadiene was added, and then, emulsion polymerization was initiated by adding 0.15 parts by weight of potassium persulfate as a polymerization initiator at a pressure of 2.8 kgf/cm² and a temperature of 41°C.

[0130] The reaction pressure at the time when the polymerization conversion rate was 40% was controlled to 2.94 kgf/cm², the reaction pressure at the time when the polymerization conversion rate was 60% was controlled to 4.62 kgf/cm², and the reaction pressure at the time when the polymerization conversion rate was 90% was controlled to 0.14 kgf/cm².

[0131] When the polymerization conversion rate reached 94%, the polymerization was terminated. Thereafter, unreacted products were removed through a deodorization process, and ammonium water, an antioxidant, and a defoaming agent were added, thereby obtaining a carboxylic acid-modified nitrile-based copolymer latex.

[0132] A latex composition for dip molding and a molded article were produced by using the obtained carboxylic acid-modified nitrile-based copolymer latex in the same manner as that of Example 1.

Example 8

[0133] 26 wt% of acrylonitrile and 4 wt% of methacrylic acid were added to a polymerization reactor installed with a stirrer and stirring was started to mix them, 0.6 parts by weight of t-dodecylmercaptan, 3.5 parts by weight of sodium dodecylbenzene sulfonate, and 120 parts by weight of water at 60°C were added, 70 wt% of isoprene was added, and then, emulsion polymerization was initiated by adding 0.3 parts by weight of potassium persulfate as a polymerization initiator at a pressure of 2.4 kgf/cm² and a temperature of 41°C.

[0134] The reaction pressure at the time when the polymerization conversion rate was 40% was controlled to 2.57 kgf/cm², the reaction pressure at the time when the polymerization conversion rate was 60% was controlled to 3.43 kgf/cm², and the reaction pressure at the time when the polymerization conversion rate was 90% was controlled to 1.32 kgf/cm².

[0135] When the polymerization conversion rate reached 94%, the polymerization was terminated. Thereafter, unreacted products were removed through a deodorization process, and ammonium water, an antioxidant, and a defoaming agent were added, thereby obtaining a carboxylic acid-modified nitrile-based copolymer latex.

[0136] A latex composition for dip molding and a molded article were produced by using the obtained carboxylic acid-modified nitrile-based copolymer latex in the same manner as that of Example 1.

Example 9

[0137] A latex composition for dip molding and a molded article were produced in the same manner as that of Example 1, except that 1 part by weight of aluminum citrate was added instead of 1.3 parts by weight of zinc oxide in the preparation of the latex composition for dip molding in Example 1.

Example 10

[0138] A latex composition for dip molding and a molded article were produced in the same manner as that of Example 4, except that 1 part by weight of aluminum citrate was added instead of 1.3 parts by weight of zinc oxide in the preparation of the latex composition for dip molding in Example 4.

Example 11

[0139] A latex composition for dip molding and a molded article were produced in the same manner as that of Example 5, except that 1 part by weight of aluminum citrate was added instead of 1.3 parts by weight of zinc oxide in the preparation of the latex composition for dip molding in Example 5.

Example 12

[0140] A latex composition for dip molding and a molded article were produced in the same manner as that of Example 6, except that 1 part by weight of aluminum citrate was added instead of 1.3 parts by weight of zinc oxide in the preparation of the latex composition for dip molding in Example 6.

Example 13

[0141] A latex composition for dip molding and a molded article were produced in the same manner as that of Example 7, except that 1 part by weight of aluminum citrate was added instead of 1.3 parts by weight of zinc oxide in the preparation of the latex composition for dip molding in Example 7.

Example 14

[0142] A latex composition for dip molding and a molded article were produced in the same manner as that of Example 1, except that 1 part by weight of sulfur, 1.0 part by weight of zinc di-n-butylthiocarbamate, and 2 parts by weight of aluminum citrate instead of 1.3 parts by weight of zinc oxide were added in the preparation of the latex composition for dip molding in Example 1.

Example 15

[0143] A latex composition for dip molding and a molded article were produced in the same manner as that of Example 4, except that 1 part by weight of sulfur, 1.0 part by weight of zinc di-n-butylthiocarbamate, and 2 parts by weight of aluminum citrate instead of 1.3 parts by weight of zinc oxide were added in Example 4.

Example 16

[0144] A latex composition for dip molding and a molded article were produced in the same manner as that of Example 5, except that 1 part by weight of sulfur, 1.0 part by weight of zinc di-n-butylthiocarbamate, and 2 parts by weight of aluminum citrate instead of 1.3 parts by weight of zinc oxide were added in the preparation of the latex composition for dip molding in Example 5.

Example 17

[0145] A latex composition for dip molding and a molded article were produced in the same manner as that of Example 6, except that 1 part by weight of sulfur, 1.0 part by weight of zinc di-n-butylthiocarbamate, and 2 parts by weight of aluminum citrate instead of 1.3 parts by weight of zinc oxide were added in the preparation of the latex composition for dip molding in Example 6.

Example 18

[0146] A latex composition for dip molding and a molded article were produced in the same manner as that of Example 7, except that 1 part by weight of sulfur, 1.0 part by weight of zinc di-n-butylthiocarbamate, and 2 parts by weight of aluminum citrate instead of 1.3 parts by weight of zinc oxide were added in the preparation of the latex composition for dip molding in Example 7.

Comparative Examples**Comparative Example 1**

[0147] 0.25 parts by weight of t-dodecylmercaptan, 4.0 parts by weight of sodium dodecylbenzene sulfonate, and 140 parts by weight of water at 60°C were added to a polymerization reactor installed with a stirrer and stirring was started to mix them, 14 wt% of acrylonitrile, 6 wt% of methacrylic acid, and 80 wt% of 1,3-butadiene were added, and emulsion polymerization was initiated by adding 0.45 parts by weight of potassium persulfate as a polymerization initiator at a pressure of 3.5 kgf/cm² and a temperature of 41°C.

[0148] The reaction pressure at the time when the polymerization conversion rate was 40% was controlled to 4.06 kgf/cm², the reaction pressure at the time when the polymerization conversion rate was 60% was controlled to 4.13 kgf/cm², and the reaction pressure at the time when the polymerization conversion rate was 90% was controlled to 3.71 kgf/cm².

[0149] When the polymerization conversion rate reached 97%, the polymerization was terminated. Thereafter, unreacted products were removed through a deodorization process, and ammonium water, an antioxidant, and a defoaming agent were added, thereby obtaining a carboxylic acid-modified nitrile-based copolymer latex.

[0150] A latex composition for dip molding and a molded article were produced by using the obtained carboxylic acid-modified nitrile-based copolymer latex in the same manner as that of Example 1.

Comparative Example 2

[0151] 25 wt% of acrylonitrile and 4.1 wt% of methacrylic acid were added to a polymerization reactor installed with a stirrer, 0.3 parts by weight of t-dodecylmercaptan, 2.5 parts by weight of sodium dodecylbenzene sulfonate, and 140 parts by weight of water at 55°C were added, 70.9 wt% of 1,3-butadiene was added, and then, emulsion polymerization

was initiated by adding 0.2 parts by weight of potassium persulfate as a polymerization initiator at a pressure of 2.9 kgf/cm² and a temperature of 40°C and starting stirring.

[0152] The reaction pressure at the time when the polymerization conversion rate was 40% was controlled to 3.34 kgf/cm², the reaction pressure at the time when the polymerization conversion rate was 60% was controlled to 3.68 kgf/cm², and the reaction pressure at the time when the polymerization conversion rate was 90% was controlled to 2.81 kgf/cm².

[0153] When the polymerization conversion rate reached 96%, the polymerization was terminated. Thereafter, unreacted products were removed through a deodorization process, and ammonium water, an antioxidant, and a defoaming agent were added, thereby obtaining a carboxylic acid-modified nitrile-based copolymer latex.

[0154] A latex composition for dip molding and a molded article were produced by using the obtained carboxylic acid-modified nitrile-based copolymer latex in the same manner as that of Example 1.

Comparative Example 3

[0155] 28 wt% of acrylonitrile and 2.5 wt% of methacrylic acid were added to a polymerization reactor installed with a stirrer, 0.5 parts by weight of t-dodecylmercaptan, 3.5 parts by weight of sodium dodecylbenzene sulfonate, and 120 parts by weight of water at 65°C were added, 69.5 wt% of 1,3-butadiene was added, and then, emulsion polymerization was initiated by adding 0.15 parts by weight of potassium persulfate as a polymerization initiator at a pressure of 1.9 kgf/cm² and a temperature of 36°C and starting stirring.

[0156] The reaction pressure at the time when the polymerization conversion rate was 40% was controlled to 2.2 kgf/cm², the reaction pressure at the time when the polymerization conversion rate was 60% was controlled to 2.7 kgf/cm², and the reaction pressure at the time when the polymerization conversion rate was 90% was controlled to 0.95 kgf/cm².

[0157] When the polymerization conversion rate reached 94%, the polymerization was terminated. Thereafter, unreacted products were removed through a deodorization process, and ammonium water, an antioxidant, and a defoaming agent were added, thereby obtaining a carboxylic acid-modified nitrile-based copolymer latex.

[0158] A latex composition for dip molding and a molded article were produced by using the obtained carboxylic acid-modified nitrile-based copolymer latex in the same manner as that of Example 1.

Comparative Example 4

[0159] 21 wt% of acrylonitrile and 2.9 wt% of methacrylic acid were added to a polymerization reactor installed with a stirrer, 0.25 parts by weight of t-dodecylmercaptan, 3.0 parts by weight of sodium dodecylbenzene sulfonate, and 110 parts by weight of water at 60°C were added, 76.1 wt% of 1,3-butadiene was added, and then, emulsion polymerization was initiated by adding 0.45 parts by weight of potassium persulfate as a polymerization initiator at a pressure of 3.0 kgf/cm² and a temperature of 41°C and starting stirring.

[0160] The reaction pressure at the time when the polymerization conversion rate was 40% was controlled to 3.15 kgf/cm², the reaction pressure at the time when the polymerization conversion rate was 60% was controlled to 3.3 kgf/cm², and the reaction pressure at the time when the polymerization conversion rate was 90% was controlled to 0.9 kgf/cm².

[0161] When the polymerization conversion rate reached 97%, the polymerization was terminated. Thereafter, unreacted products were removed through a deodorization process, and ammonium water, an antioxidant, and a defoaming agent were added, thereby obtaining a carboxylic acid-modified nitrile-based copolymer latex.

[0162] A latex composition for dip molding and a molded article were produced by using the obtained carboxylic acid-modified nitrile-based copolymer latex in the same manner as that of Example 1.

Comparative Example 5

[0163] 18 wt% of acrylonitrile and 4.5 wt% of methacrylic acid were added to a polymerization reactor installed with a stirrer, 0.6 parts by weight of t-dodecylmercaptan, 3.5 parts by weight of sodium dodecylbenzene sulfonate, and 110 parts by weight of water at 75°C were added, 77.5 wt% of 1,3-butadiene was added, and then, emulsion polymerization was initiated by adding 0.3 parts by weight of potassium persulfate as a polymerization initiator at a pressure of 3.3 kgf/cm² and a temperature of 41°C and starting stirring.

[0164] The reaction pressure at the time when the polymerization conversion rate was 40% was controlled to 3.53 kgf/cm², the reaction pressure at the time when the polymerization conversion rate was 60% was controlled to 4.69 kgf/cm², and the reaction pressure at the time when the polymerization conversion rate was 90% was controlled to 3.5 kgf/cm².

[0165] When the polymerization conversion rate reached 95%, the polymerization was terminated. Thereafter, unre-

acted products were removed through a deodorization process, and ammonium water, an antioxidant, and a defoaming agent were added, thereby obtaining a carboxylic acid-modified nitrile-based copolymer latex.

[0166] A latex composition for dip molding and a molded article were produced by using the obtained carboxylic acid-modified nitrile-based copolymer latex in the same manner as that of Example 1.

Comparative Example 6

[0167] 0.25 parts by weight of t-dodecylmercaptan, 3.0 parts by weight of sodium dodecylbenzene sulfonate, and 110 parts by weight of water at 60°C were added to a polymerization reactor installed with a stirrer and stirring was started to mix them, 31 wt% of acrylonitrile and 6 wt% of methacrylic acid were added, 63 wt% of 1,3-butadiene were added, and emulsion polymerization was initiated by adding 0.45 parts by weight of potassium persulfate as a polymerization initiator at a pressure of 2.0 kgf/cm² and a temperature of 36°C and starting stirring.

[0168] The reaction pressure at the time when the polymerization conversion rate was 40% was controlled to 2.2 kgf/cm², the reaction pressure at the time when the polymerization conversion rate was 60% was controlled to 2.6 kgf/cm², and the reaction pressure at the time when the polymerization conversion rate was 90% was controlled to 1.8 kgf/cm².

[0169] When the polymerization conversion rate reached 97%, the polymerization was terminated. Thereafter, unreacted products were removed through a deodorization process, and ammonium water, an antioxidant, and a defoaming agent were added, thereby obtaining a carboxylic acid-modified nitrile-based copolymer latex.

[0170] A latex composition for dip molding and a molded article were produced by using the obtained carboxylic acid-modified nitrile-based copolymer latex in the same manner as that of Example 1.

Comparative Example 7

[0171] A latex composition for dip molding and a molded article were produced in the same manner as that of Comparative Example 1, except that 1 part by weight of aluminum citrate was added instead of 1.3 parts by weight of zinc oxide in the preparation of the latex composition for dip molding in Comparative Example 1.

Comparative Example 8

[0172] A latex composition for dip molding and a molded article were produced in the same manner as that of Comparative Example 2, except that 1 part by weight of aluminum citrate was added instead of 1.3 parts by weight of zinc oxide in the preparation of the latex composition for dip molding in Comparative Example 2.

Comparative Example 9

[0173] A latex composition for dip molding and a molded article were produced in the same manner as that of Comparative Example 6, except that 1 part by weight of aluminum citrate was added instead of 1.3 parts by weight of zinc oxide in the preparation of the latex composition for dip molding in Comparative Example 6.

Comparative Example 10

[0174] A latex composition for dip molding and a molded article were produced in the same manner as that of Comparative Example 1, except that 1 part by weight of sulfur, 1.0 part by weight of zinc di-n-butyldithiocarbamate, and 2 parts by weight of aluminum citrate instead of 1.3 parts by weight of zinc oxide were added in the preparation of the latex composition for dip molding in Comparative Example 1.

Comparative Example 11

[0175] A latex composition for dip molding and a molded article were produced in the same manner as that of Comparative Example 2, except that 1 part by weight of sulfur, 1.0 part by weight of zinc di-n-butyldithiocarbamate, and 2 parts by weight of aluminum citrate instead of 1.3 parts by weight of zinc oxide were added in the preparation of the latex composition for dip molding in Comparative Example 2.

Comparative Example 12

[0176] A latex composition for dip molding and a molded article were produced in the same manner as that of Comparative Example 6, except that 1 part by weight of sulfur, 1.0 part by weight of zinc di-n-butyldithiocarbamate, and 2

parts by weight of aluminum citrate instead of 1.3 parts by weight of zinc oxide were added in the preparation of the latex composition for dip molding in Comparative Example 6.

Experimental Examples

Experimental Example 1

[0177] Physical properties of the carboxylic acid-modified nitrile-based copolymer latex prepared in each of Examples 1 to 8 and Comparative Examples 1 to 6 were measured by the following method. The results are shown in Table 1.

(1) CV_0 (mm^2/s): A pH of each carboxylic acid-modified nitrile-based copolymer latex having a solid content of 44% to 47% was adjusted to 8.8 to 9.1 using 10% ammonium water, and then, the carboxylic acid-modified nitrile-based copolymer latex was dissolved in methyl ethyl ketone (MEK) at a concentration of 2.55 wt% to be dispersed well. Thereafter, 10 ml of the carboxylic acid-modified nitrile-based copolymer latex was added to a Cannon-Fenske routine type capillary viscometer (SI Analytics GmbH Type No. 520 13), the time for the copolymer latex to pass through a capillary at 25°C was measured, and a viscosity was calculated using the following Equation 2.

[Equation 2]

$$CV_0 = K \times t$$

In Equation 2, K is a constant (mm^2/s^2) of the capillary, and t is a time (s) for the copolymer latex to pass through the capillary.

(2) CV_D (mm^2/s), P: A pH of each carboxylic acid-modified nitrile-based copolymer latex having a solid content of 44% to 47% was adjusted to 8.8 to 9.1 using 10% ammonium water, and then, the carboxylic acid-modified nitrile-based copolymer latex was dissolved in methyl ethyl ketone (MEK) at a concentration of 2.55 wt% to be dispersed well. Thereafter, de-swelling was performed by applying an energy of 55 kcal to 65 kcal using the ultrasonicator (Bransonic® M Mechanical Bath 5800) for 40 minutes, 10 ml of the carboxylic acid-modified nitrile-based copolymer latex was added to the Cannon-Fenske routine type capillary viscometer (SI Analytics GmbH Type No. 520 13), the time for the copolymer latex to pass through the capillary at 25°C was measured, and a viscosity (CV_D) was calculated using Equation 2. P was measured by dividing the calculated CV_D by CV_0 ($P = CV_D/CV_0$).

(3) M: A pH of each carboxylic acid-modified nitrile-based copolymer latex having a solid content of 44% to 47% was adjusted to 8.8 to 9.1 using 10% ammonium water, and the copolymer latex was applied onto a substrate using a cast method or the like, drying was performed at 130°C to 140°C for 40 minutes to obtain a dried film, and a weight (f_1) of the dried film was measured. Subsequently, the dried film was dipped into methyl ethyl ketone at 25°C for 48 hours, a weight (f_2) of the dipped film was measured, and drying was performed at 170°C for 30 minutes to remove the methyl ethyl ketone. Then, a weight (f_3) of the film after removing the methyl ethyl ketone was measured. An insolubility (m_1) and a swelling index (m_2) of the film in the methyl ethyl ketone solvent were measured from the measurement results of these weights according to the following Equations 3 and 4, and M was measured according to Equation 5.

[Equation 3]

Insolubility in methyl ethyl ketone solvent (m_1) =

$$f_3/f_1$$

[Equation 4]

Swelling index in methyl ethyl ketone solvent (m_2) =

$$(f_2 - f_3)/f_3$$

[Equation 5]

$$M = m_1 \times m_2$$

(4) pKa: The carboxylic acid-modified nitrile-based copolymer latex according to Example 1 was diluted using deionized water to a solid content of 10%, a pH thereof was increased to 12 using a 1.25% aqueous potassium hydroxide solution, and stirring was performed at 50°C for 2 hours to remove ammonia in the aqueous solution. The obtained diluted solution was cooled to room temperature, a pH of the solution was decreased to 2 or less using a 1% diluted aqueous hydrochloric acid solution, and the solution was stirred at a temperature of 50°C for 2 hours. Next, carbon dioxide in the aqueous solution was removed, the obtained diluted solution was cooled to room temperature, and an acid-base titration graph was obtained using an aqueous potassium hydroxide solution having a concentration of exactly 3%. The results are shown in Tables 1 and 2. FIG. 1 is a graph showing a change in pH according to an addition amount of KOH. The amount of carboxylic acid calculated by the addition amount of KOH between the first inflection point and the second inflection point in FIG. 1 is the amount of acid present on a surface. FIG. 2 is a derivative curve of the addition amount of KOH according to the pH. The vertex of the quadratic function is a pKa of the carboxylic acid-modified nitrile-based copolymer latex. In addition, a pKa of the remaining carboxylic acid-modified nitrile-based copolymer latex according to each of Examples 2 to 8 and Comparative Examples 1 to 6 was confirmed in the same manner.

[Table 1]

	CV ₀ (mm ² /s)	CV _D (mm ² /s)	P	M	pKa
Example 1	2.13	1.96	0.92	10.7	9.93
Example 2	1.26	1.21	0.96	13.2	9.68
Example 3	2.81	2.61	0.93	7.0	10.12
Example 4	2.57	2.31	0.90	9.1	10.07
Example 5	1.65	1.62	0.98	11.7	9.83
Example 6	3.00	2.73	0.91	10.3	9.84
Example 7	1.00	0.96	0.96	9.8	9.92
Example 8	2.68	2.49	0.93	9.9	9.95
Comparative Example 1	0.40	0.30	0.76	20.2	9.27
Comparative Example 2	2.57	2.03	0.79	11.3	9.85
Comparative Example 3	2.81	2.36	0.84	9.3	9.97
Comparative Example 4	1.65	1.34	0.81	9.5	9.97
Comparative Example 5	1.26	1.01	0.80	14.4	9.55
Comparative Example 6	5.50	5.01	0.91	20.2	9.28

[0178] Referring to Table 1, it could be confirmed that in the case of Examples 1 to 8, physical properties of the latex itself were improved through optimization of the structure of the latex particles by adjusting the entanglement degree or the branch degree and the molecular weight of the carboxylic acid-modified nitrile-based copolymer and the acid distribution of the copolymer in the latex particles in the polymerization step by controlling process factors for the carboxylic acid-modified nitrile-based copolymer latex according to the present invention, and thus, CV₀ was in a range of 1.0 to 3.0, and the value of P was in a range of 0.85 to 1.

[0179] On the other hand, in the case of Comparative Examples 1 to 6, the structure of the latex particles was not optimized as in the present invention, and thus, the physical properties of the latex were not improved.

Experimental Example 2

[0180] Physical properties of the molded article produced in each of Examples 1 to 8 and Comparative Examples 1 to 6 were measured by the following method. The results are shown in Table 2.

(1) Modulus at 500% (MPa): A dumbbell-shaped specimen was prepared from the molded article according to the ASTM D-412 method, the specimen was pulled at a cross-head speed of 500 mm/min using a measurement device (U.T.M.) (3345 model, Instron), and a stress when elongation was 500% was measured.

(2) Stress retention rate (%): A dumbbell-shaped specimen was prepared from the molded article according to the ASTM D-412 method, the specimen was pulled at a cross-head speed of 300 mm/min using a measurement device (U.T.M.) (3345 model, Instron) until elongation calculated according to the following Equation 6 reached 100%, a stress reduction

for 5 minutes was measured, and a stress retention rate was calculated according to the following Equation 7.

[Equation 6]

$$\text{Elongation (\%)} = (\text{length after specimen elongation} / \text{initial length of specimen}) \times 100$$

[Equation 7]

$$\text{Stress retention rate (\%)} = (\text{load value 5 minutes after specimen elongation} / \text{initial load value of specimen elongation}) \times 100$$

[Table 2]

	Cross-linking agent composition (parts by weight)					Modulus at 500% (MPa)	Stress retention rate (%)
	S	ZDBC	ZnO	Al	TiO ₂		
Example 1	1	1	1.3	0	1	16.1	47
Example 2	1	1	1.3	0	1	11.6	50.1
Example 3	1	1	1.3	0	1	15.3	46.6
Example 4	1	1	1.3	0	1	18.5	45
Example 5	1	1	1.3	0	1	10.4	51.9
Example 6	1	1	1.3	0	1	18.1	45.4
Example 7	1	1	1.3	0	1	9.7	49.2
Example 8	1	1	1.3	0	1	16.9	47
Comparative Example 1	1	1	1.3	0	1	33.6	33
Comparative Example 2	1	1	1.3	0	1	22.4	34.4
Comparative Example 3	1	1	1.3	0	1	23.1	39.4
Comparative Example 4	1	1	1.3	0	1	22.3	37
Comparative Example 5	1	1	1.3	0	1	21.4	34.9
Comparative Example 6	1	1	1.3	0	1	-	45.4

[0181] In Table 2, S represents sulfur, ZDBC represents di-n-butylthiocarbamate, ZnO represents zinc oxide, Al represents aluminum citrate, and TiO₂ represents titanium oxide.

[0182] Referring to Table 2, it could be confirmed that in the case of Examples 1 to 8, in the molded article formed of the latex composition for dip molding comprising the carboxylic acid-modified nitrile-based copolymer latex according to the present invention, the stress retention rate was high and the value of the modulus at 500% was small, which showed that both the elasticity and softness were improved.

[0183] On the other hand, it could be confirmed that in the case of Comparative Examples 1 to 6, in the molded article formed of the latex composition for dip molding comprising the unimproved carboxylic acid-modified nitrile-based copolymer latex unlike in the present invention, the stress retention rate and the value of the modulus at 500% were significantly reduced as compared to those in Examples.

[0184] In addition, in the case of Comparative Example 6, in the molded article formed of the latex composition for dip molding comprising the unimproved carboxylic acid-modified nitrile-based copolymer latex unlike in the present invention, the measured elongation was less than 500%, and thus, the value of the modulus at 500% could not be measured.

Experimental Example 3

[0185] In order to show a change of the physical properties of the molded article according to the change of the cross-linking agent, the physical properties of the molded article produced in each of Examples 9 to 18 and Comparative Examples 7 to 12 were measured in the same manner as that of Experimental Example 2. The results are shown in Table 3.

[Table 3]

		S	ZDBC	ZnO	Type of latex	Cross-linking agent composition (parts by weight)	Modulus at 500% (MPa)	Stress retention rate
					Al	TiO ₂		(%)
Example 9	Example 1	1	1	0	1	1	9.4	54.9
Example 10	Example 4	1	1	0	1	1	10.8	54.4
Example 11	Example 5	1	1	0	1	1	6.1	56.9
Example 12	Example 6	1	1	0	1	1	13.9	54.7
Example 13	Example 7	1	1	0	1	1	6.4	56
Example 14	Example 1	0	0	0	2	1	10.1	54.8
Example 15	Example 4	0	0	0	2	1	11.6	53.9
Example 16	Example 5	0	0	0	2	1	6.6	57.2
Example 17	Example 6	0	0	0	2	1	13.1	54
Example 18	Example 7	0	0	0	2	1	7.2	56.6
Comparative Example 7	Comparative Example 1	1	1	0	1	1	20.6	41.2
Comparative Example 8	Comparative Example 2	1	1	0	1	1	15.8	43.5
Comparative Example 9	Comparative Example 6	1	1	0	1	1	17.4	54.8
Comparative Example 10	Comparative Example 1	0	0	0	2	1	21.9	40.6
Comparative Example 11	Comparative Example 2	0	0	0	2	1	16.3	43.4
Comparative Example 12	Comparative Example 6	0	0	0	2	1	18.4	54.4

[0186] Referring to Table 3, it could be confirmed that in the case of Examples 9 to 18, in the molded article formed of the latex composition for dip molding comprising the carboxylic acid-modified nitrile-based copolymer latex according to the present invention, even in a case where the cross-linking agent composition was changed, both the modulus at 500% and the stress retention rate were improved.

[0187] On the other hand, it could be confirmed that in the case of Comparative Examples 7 to 12, in the molded article formed of the latex composition for dip molding comprising the unimproved carboxylic acid-modified nitrile-based copolymer latex unlike in the present invention, even in a case where the cross-linking agent composition was changed, the

stress retention rate and the value of the modulus at 500% were significantly reduced as compared to those in Examples.

Claims

1. A latex composition for dip molding comprising: a carboxylic acid-modified nitrile-based copolymer latex, satisfying the following General Formula 1 and General Formula 2,

wherein the carboxylic acid-modified nitrile-based copolymer includes a conjugated diene-based monomer-derived unit, an ethylenic unsaturated nitrile-based monomer-derived unit, and an ethylenic unsaturated acid monomer-derived unit:

[General Formula 1]

$$1.0 \text{ mm}^2/\text{s} \leq CV_0 \leq 3.0 \text{ mm}^2/\text{s}$$

[General Formula 2]

$$0.85 \leq P \leq 1.0, P = CV_D/CV_0$$

in General Formula 1 and General Formula 2, CV_0 represents a capillary viscosity of the carboxylic acid-modified nitrile-based copolymer latex measured in a swollen state in a methyl ethyl ketone solvent, and CV_D represents a capillary viscosity of the carboxylic acid-modified nitrile-based copolymer latex in a de-swollen state.

2. The latex composition for dip molding of claim 1, wherein the conjugated diene-based monomer includes one or more selected from the group consisting of 1,3-butadiene, 2,3-dimethyl-1,3-butadiene, 2-ethyl-1,3-butadiene, 1,3-pentadiene, and isoprene.
3. The latex composition for dip molding of claim 1, wherein the ethylenic unsaturated nitrile-based monomer includes one or more selected from the group consisting of acrylonitrile, methacrylonitrile, fumaronitrile, α -chloronitrile, and α -cyanoethylacrylonitrile.
4. The latex composition for dip molding of claim 1, wherein the ethylenic unsaturated acid monomer includes one or more selected from the group consisting of acrylic acid, methacrylic acid, itaconic acid, maleic acid, fumaric acid, maleic acid anhydride, citraconic acid anhydride, styrenesulfonic acid, monobutyl fumarate, monobutyl maleate, and mono-2-hydroxypropyl maleate.
5. The latex composition for dip molding of claim 1, wherein the conjugated diene-based monomer is 1,3-butadiene or isoprene,
the ethylenic unsaturated nitrile-based monomer is acrylonitrile, and
the ethylenic unsaturated acid monomer is methacrylic acid.
6. The latex composition for dip molding of claim 1, wherein the carboxylic acid-modified nitrile-based copolymer further includes an ethylenic unsaturated monomer-derived unit.
7. The latex composition for dip molding of claim 1, wherein the carboxylic acid-modified nitrile-based copolymer includes 15 wt% to 30 wt% of the ethylenic unsaturated nitrile-based monomer-derived unit, 1.5 wt% to 5.5 wt% of the ethylenic unsaturated acid monomer-derived unit, and 64.5 wt% to 83.5 wt% of the conjugated diene-based monomer-derived unit.
8. The latex composition for dip molding of claim 1, wherein P is 0.9 to 1.
9. The latex composition for dip molding of claim 1, wherein the carboxylic acid-modified nitrile-based copolymer latex satisfies the following General Formula 3 and General Formula 4:

[General Formula 3]

$$5 \leq M \leq 20, M = m_1 \times m_2$$

[General Formula 4]

$$9.2 \leq pK_a \leq 10.5$$

in General Formula 3, m_1 represents an insolubility of a dried film of the carboxylic acid-modified nitrile-based copolymer latex in the methyl ethyl ketone solvent, and m_2 represents a swelling index of the dried film of the carboxylic acid-modified nitrile-based copolymer latex in the methyl ethyl ketone solvent, and in General Formula 4, a pK_a represents a degree of ionization of an ethylenic unsaturated acid of the carboxylic acid-modified nitrile-based copolymer latex.

10. The latex composition for dip molding of claim 9, wherein the pK_a is 9.5 to 10.2.
11. The latex composition for dip molding of claim 1, further comprising a cross-linking agent composition.
12. The latex composition for dip molding of claim 11, wherein the cross-linking agent composition includes one or more selected from the group consisting of a vulcanizing agent, a vulcanization accelerator, a divalent cationic metal oxide, and a polyvalent metal cationic compound.
13. The latex composition for dip molding of claim 12, wherein the cross-linking agent composition includes a vulcanizing agent, a vulcanization accelerator, and a polyvalent metal cationic compound.
14. The latex composition for dip molding of claim 12, wherein the cross-linking agent composition includes a polyvalent metal cationic compound.
15. A molded article comprising a layer derived from the latex composition for dip molding of claim 1.
16. The molded article of claim 15, wherein the molded article has a modulus at 500% of 4 to 20.5 and a stress retention rate of 40% to 70%.

FIG. 1

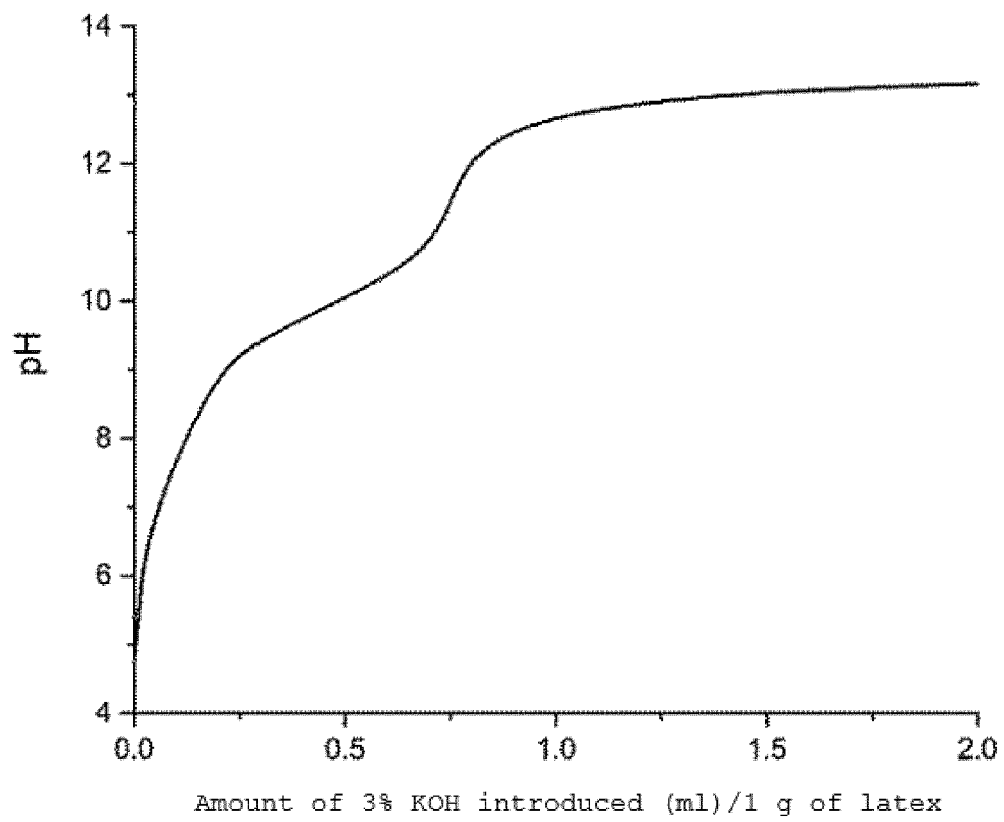
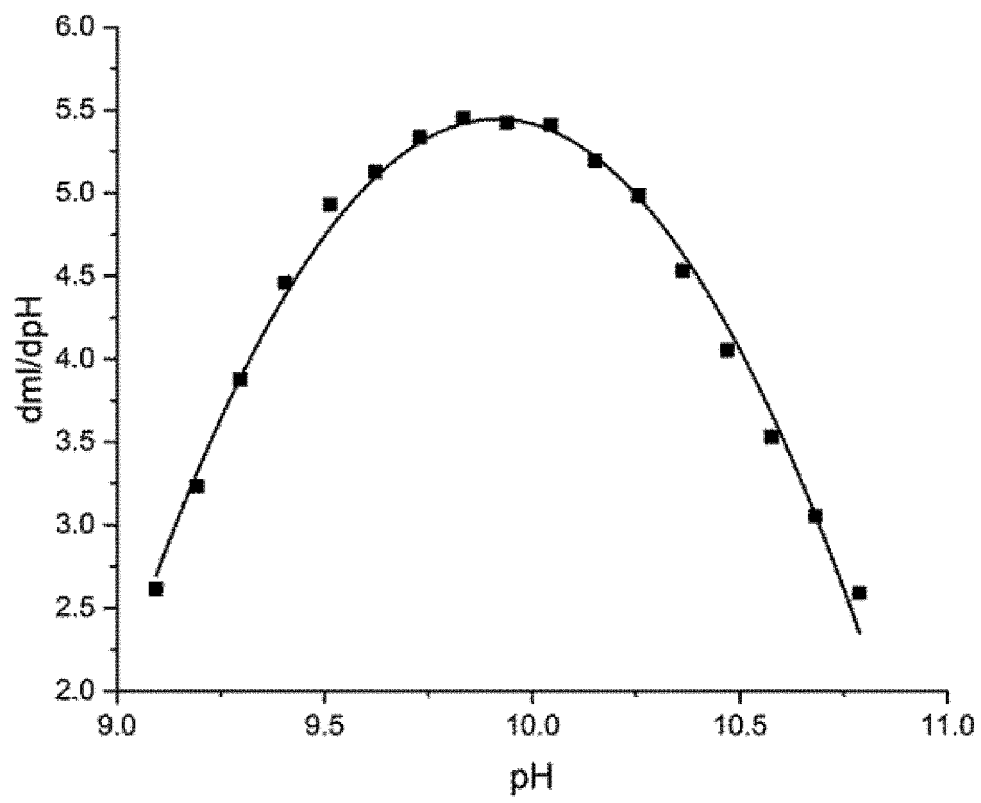


FIG. 2



INTERNATIONAL SEARCH REPORT

International application No.

PCT/KR2021/001626

A. CLASSIFICATION OF SUBJECT MATTER C08L 13/02(2006.01)i; C08L 9/04(2006.01)i; C08J 3/02(2006.01)i; C08F 236/12(2006.01)i; C08K 3/011(2018.01)i; C08K 3/22(2006.01)i; C08J 5/02(2006.01)i; B29C 41/14(2006.01)i According to International Patent Classification (IPC) or to both national classification and IPC																		
B. FIELDS SEARCHED Minimum documentation searched (classification system followed by classification symbols) C08L 13/02(2006.01); B29C 41/14(2006.01); C08F 2/22(2006.01); C08F 2/38(2006.01); C08F 236/06(2006.01); C08F 236/12(2006.01); C08L 9/04(2006.01) Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched Korean utility models and applications for utility models: IPC as above Japanese utility models and applications for utility models: IPC as above Electronic data base consulted during the international search (name of data base and, where practicable, search terms used) eKOMPASS (KIPO internal) & keywords: 라텍스(latex), 압력(pressure), 부타디엔(butadiene), 아크릴로니트릴(acrylonitrile), 메타크릴산(methacrylic acid)																		
C. DOCUMENTS CONSIDERED TO BE RELEVANT <table border="1"> <thead> <tr> <th>Category*</th> <th>Citation of document, with indication, where appropriate, of the relevant passages</th> <th>Relevant to claim No.</th> </tr> </thead> <tbody> <tr> <td>A</td> <td>US 2005-0171272 A1 (OTA, H. et al.) 04 August 2005 (2005-08-04) See claim 1; manufacturing example 2; and table 1.</td> <td>1-16</td> </tr> <tr> <td>A</td> <td>JP 2014-088536 A (JINTEX CORPORATION LTD.) 15 May 2014 (2014-05-15) See claims 1-3 and 5.</td> <td>1-16</td> </tr> <tr> <td>A</td> <td>US 2020-0062879 A1 (SYNTHOMER SDN. BHD.) 27 February 2020 (2020-02-27) See entire document.</td> <td>1-16</td> </tr> <tr> <td>A</td> <td>KR 10-1811343 B1 (KOREA KUMHO PETROCHEMICAL CO., LTD.) 26 December 2017 (2017-12-26) See entire document.</td> <td>1-16</td> </tr> <tr> <td>A</td> <td>US 2017-0327669 A1 (SYNTHOMER SDN. BHD.) 16 November 2017 (2017-11-16) See entire document.</td> <td>1-16</td> </tr> </tbody> </table>	Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.	A	US 2005-0171272 A1 (OTA, H. et al.) 04 August 2005 (2005-08-04) See claim 1; manufacturing example 2; and table 1.	1-16	A	JP 2014-088536 A (JINTEX CORPORATION LTD.) 15 May 2014 (2014-05-15) See claims 1-3 and 5.	1-16	A	US 2020-0062879 A1 (SYNTHOMER SDN. BHD.) 27 February 2020 (2020-02-27) See entire document.	1-16	A	KR 10-1811343 B1 (KOREA KUMHO PETROCHEMICAL CO., LTD.) 26 December 2017 (2017-12-26) See entire document.	1-16	A	US 2017-0327669 A1 (SYNTHOMER SDN. BHD.) 16 November 2017 (2017-11-16) See entire document.	1-16
Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.																
A	US 2005-0171272 A1 (OTA, H. et al.) 04 August 2005 (2005-08-04) See claim 1; manufacturing example 2; and table 1.	1-16																
A	JP 2014-088536 A (JINTEX CORPORATION LTD.) 15 May 2014 (2014-05-15) See claims 1-3 and 5.	1-16																
A	US 2020-0062879 A1 (SYNTHOMER SDN. BHD.) 27 February 2020 (2020-02-27) See entire document.	1-16																
A	KR 10-1811343 B1 (KOREA KUMHO PETROCHEMICAL CO., LTD.) 26 December 2017 (2017-12-26) See entire document.	1-16																
A	US 2017-0327669 A1 (SYNTHOMER SDN. BHD.) 16 November 2017 (2017-11-16) See entire document.	1-16																
☐ Further documents are listed in the continuation of Box C. ☒ See patent family annex.																		
<table border="0"> <tr> <td style="vertical-align: top;"> * Special categories of cited documents: "A" document defining the general state of the art which is not considered to be of particular relevance "D" document cited by the applicant in the international application "E" earlier application or patent but published on or after the international filing date "L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified) "O" document referring to an oral disclosure, use, exhibition or other means "P" document published prior to the international filing date but later than the priority date claimed </td> <td style="vertical-align: top;"> "T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention "X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone "Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art "&" document member of the same patent family </td> </tr> </table>	* Special categories of cited documents: "A" document defining the general state of the art which is not considered to be of particular relevance "D" document cited by the applicant in the international application "E" earlier application or patent but published on or after the international filing date "L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified) "O" document referring to an oral disclosure, use, exhibition or other means "P" document published prior to the international filing date but later than the priority date claimed	"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention "X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone "Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art "&" document member of the same patent family																
* Special categories of cited documents: "A" document defining the general state of the art which is not considered to be of particular relevance "D" document cited by the applicant in the international application "E" earlier application or patent but published on or after the international filing date "L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified) "O" document referring to an oral disclosure, use, exhibition or other means "P" document published prior to the international filing date but later than the priority date claimed	"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention "X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone "Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art "&" document member of the same patent family																	
Date of the actual completion of the international search 27 May 2021	Date of mailing of the international search report 27 May 2021																	
Name and mailing address of the ISA/KR Korean Intellectual Property Office Government Complex-Daejeon Building 4, 189 Cheongsaro, Seo-gu, Daejeon 35208 Facsimile No. +82-42-481-8578	Authorized officer Telephone No.																	

Form PCT/ISA/210 (second sheet) (July 2019)

INTERNATIONAL SEARCH REPORT
Information on patent family members

International application No.

PCT/KR2021/001626

Patent document cited in search report	Publication date (day/month/year)	Patent family member(s)	Publication date (day/month/year)
US 2005-0171272 A1	04 August 2005	CN 1297602 C	31 January 2007
		CN 1656166 A	17 August 2005
		EP 1487915 A1	22 December 2004
		JP 2003-277523 A	02 October 2003
		JP 3852356 B2	29 November 2006
		MY 144286 A	29 August 2011
		US 2007-0083013 A1	12 April 2007
		WO 03-080722 A1	02 October 2003
JP 2014-088536 A	15 May 2014	CN 103788298 A	14 May 2014
		JP 5719388 B2	20 May 2015
		TW 201416381 A	01 May 2014
		TW 1481629 B	21 April 2015
US 2020-0062879 A1	27 February 2020	BR 112019010268 A2	10 September 2019
		CN 110072936 A	30 July 2019
		EP 3555198 A1	23 October 2019
		JP 2020-503401 A	30 January 2020
		KR 10-2019-0090391 A	01 August 2019
		TW 201835194 A	01 October 2018
		WO 2018-111087 A1	21 June 2018
KR 10-1811343 B1	26 December 2017	KR 10-2017-0110956 A	12 October 2017
US 2017-0327669 A1	16 November 2017	AR 103143 A1	19 April 2017
		CN 107636032 A	26 January 2018
		CN 107636032 B	07 April 2020
		EP 3230361 A1	18 October 2017
		EP 3230361 B1	28 November 2018
		JP 2018-505916 A	01 March 2018
		JP 6431981 B2	28 November 2018
		KR 10-2017-0094156 A	17 August 2017
		TW 201629106 A	16 August 2016
		US 10377882 B2	13 August 2019
		UY 36430 A	31 August 2016
		WO 2016-093689 A1	16 June 2016

REFERENCES CITED IN THE DESCRIPTION

This list of references cited by the applicant is for the reader's convenience only. It does not form part of the European patent document. Even though great care has been taken in compiling the references, errors or omissions cannot be excluded and the EPO disclaims all liability in this regard.

Patent documents cited in the description

- KR 1020200039060 [0001]
- KR 1020200039067 [0001]
- KR 1020200039076 [0001]
- KR 1020200039074 [0001]
- KR 1020200039065 [0001]
- KR 1020200039071 [0001]
- KR 1020200172992 [0001]